

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:06:39 ; Search time 15 Seconds

(without alignments)
1769.504 Million cell updates/sec

Title: US-10-092-404-2

Perfect score: 1520

Sequence: 1 RLRSLSLHFLFMGASEQDL.....RYTCQVEHPGLDPLIVWE 276

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 283308 seqs, 96168682 residues

Total number of hits satisfying chosen parameters: 283308

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

PIR 76:**

1: Pirl:**

2: Pirl:**

3: Pirl:**

4: Pirl:**

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match %	Length	DB ID	Description
1	1140	75.0	359	2 JCS382	hereditary hemochr
2	542.5	35.7	341	2 A57136	class I histocompa
3	523	34.4	361	1 HLRB	MHC class I histoc
4	523	34.4	361	2 I46858	MHC class I RLA pr
5	520	34.2	332	2 S06424	MHC class I histoc
6	517	34.0	365	2 I36961	MHC class I protei
7	516	33.9	361	2 B27638	MHC class I histoc
8	515	33.9	365	2 I83063	A11.2 - human
9	514	33.8	365	2 A47636	MHC class I histoc
10	514	33.8	365	2 I56039	HLA-A30.3 precuro
11	512	33.7	370	1 HLHUA3	MHC class I histoc
12	510	33.6	365	2 I38439	MHC class I histoc
13	509	33.5	365	2 I37542	MHC class I histoc
14	509	33.5	365	2 I38442	gene HLA-A-0205 pr
15	509	33.5	365	2 I61902	MHC class I histoc
16	508	33.4	365	2 I72170	MHC class I histoc
17	508	33.4	365	2 I38441	gene HLA-A-6802 pr
18	506	33.3	365	1 HLHUA2	MHC class I histoc
19	506	33.3	365	2 I37482	MHC class I histoc
20	506	33.3	365	2 I38519	MHC class I histoc
21	506	33.3	365	2 I84448	MHC class I histoc
22	505	33.2	365	2 I38610	MHC class I histoc
23	505	33.2	365	2 I37470	HLA-A*0210 - human
24	504	33.2	365	2 T28149	MHC class I histoc
25	504	33.2	364	2 S03535	class I histocompa
26	503	33.1	365	2 I37476	MHC class I histoc
27	503	33.1	365	2 I37478	MHC class I histoc
28	503	33.1	365	2 I38443	gene HLA-A-0203 pr
29	503	33.1	365	2 I61857	MHC HLA-A2.4a chai

RESULT 1

JCS382

hereditary hemochromatosis protein precursor - mouse

C:Species: Mus musculus (house mouse)

C:Date: 02-Jun-1997 #sequence_revision 18-Jul-1997 #text_change 05-Nov-1999

C:Accession: JCS382

R:Hashimoto, K.; Hirai, M.; Kurosawa, Y.

Biochem. Biophys. Res. Commun. 230, 35-39, 1997

A:Title: Identification of a mouse homolog for the human hereditary haemochromatosis ca

A:Reference number: JCS382; MUID:97148566; PMID:9020055

A:Accession: JCS382

A:Status: nucleic acid sequence not shown

A:Molecule type: DNA

A:Residues: 1-359 <HAS>

A:Cross-references: GB:U66849; NID:gl519484; PIDN:AAB07525.1; PID:gl519485

C:Comment: This protein plays a role in iron metabolism.

C:Genetics:

A:Gene: mr2

C:Superfamily: class I histocompatibility antigen; immunoglobulin homology

F:1-29/Domain: signal sequence #status predicted <SIG>

F:30-359/Product: hereditary haemochromatosis protein #status predicted <MAT>

F:30-117/Domain: alpha 1 #status predicted <ALF1>

F:118-217/Domain: alpha 2 #status predicted <ALF2>

F:218-309/Domain: alpha 3 #status predicted <ALF3>

F:314-340/Domain: transmembrane #status predicted <TRM>

F:341-359/Domain: intracellular #status predicted <INT>

Query Match 75.0%; Score 1140; DB 2; Length 359;

Best Local Similarity 72.2%; Pred. No. 2.3e-87;

Matches 203; Conservative 30; Mismatches 40; Indels 8; Gaps 1;

QY	4	RSLSLHFLFMGASEODLGLSFLAAGYDDQLFVYDDSESRVETPTWVSRISQMWL	63
DB	30	RSLSLHFLFMGASEPDGLPLFARGYDDQLFVSYNHSRAEPAPWILEQTSSQLWL	89
QY	64	QLSLSLHFLFMGASEODLGLSFLAAGYDDQLFVYDDSESRVETPTWVSRISQMWL	115
DB	90	HLSQLKGDYMFVDFWTIMGNVNSKVTGLGVVSESHILQVLGCEVHEDNSTSGFWR	149
QY	116	GYDGDHLEFCDDTLDWRAAPRAWPTKLEWERIKIRARQNYLERDPCPAQLQLLEL	175
DB	150	GYDGDHLEFCDDTLDWRAAPRAWPTKLEWERIKIRARQNYLERDPCPAQLQLLEL	209
QY	176	GRGVLDQVPLVAVKVTHTVTSVTLRCALNYPQNTIMKWLKDKQPMDAKEFPKDV	235
DB	210	GRGVLDQVPLVAVKVTHTVTSVTLRCALNYPQNTIMKWLKDKQPMDAKEFPKDV	269
QY	236	PNGDGTQGWITLAVPPGEQRYTCQVEHPGLDPLIVWE	276
DB	270	PNGDGTQGWITLAVAPGDETRFTCQVEHPGLDPLIVWE	310

ALIGNMENTS

30	502.5	33.1	341	2	JCS5663	major histocompati
31	502	33.0	357	2	I36965	MHC class I protei
32	501.5	33.0	362	2	A45845	MHC class I histoc
33	501	33.0	365	2	I61856	MHC class I histoc
34	501	33.0	365	2	I54493	MHC class I histoc
35	500	32.9	273	1	HLHU69	MHC class I histoc
36	500	32.9	365	2	S77963	MHC class I histoc
37	500	32.9	365	2	S01171	class I histocompa
38	500	32.9	365	2	I54416	HLA-AW4 protein -
39	499	32.8	365	2	I37483	HLA-AW34.2 antigen
40	498	32.8	273	1	HLHUAW	MHC class I histoc
41	498	32.8	360	2	A27638	MHC class I histoc
42	498	32.8	365	2	I72171	HLA-AW33.1, HLA-AW
43	497.5	32.7	339	2	I56071	MHC class I histoc
44	497	32.7	279	2	JX0353	zinc-alpha 2-glyco
45	497	32.7	362	2	I68724	MHC class I histoc

RESULT 2

A57136
Class I histocompatibility antigen related protein MR1 precursor - human
C:Species: Homo sapiens (man)
C:Date: 23-Feb-1996 #sequence_revision 23-Feb-1996 #text_change 23-Jul-1999
C:Accession: A57136
R:Hashimoto, K.; Hirai, M.; Kurosawa, Y.
Science 269, 693-695, 1995
A:Title: A gene outside the human MHC related to classical HLA class I genes.
A:Reference number: A57136; MUID:95350662; PMID:7624800
A:Accession: A57136
A:Status: preliminary
A:Molecule type: mRNA
A:Residues: 1-341 <HAS>
A:Cross-references: GB:U22963; NID:9940353; PIDN:AAC50174.1; PID:g940354
C:Genetics:
A:Gene: GDB:HLALS
A:Cross-references: GDB:683188; OMIM:600764
A:Map position: 1q25.3-1q25.3
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology

Query Match 35.78; Score 542.5; DB 2; Length 341;
Best Local Similarity 39.5%; Pred. No. 1.2e-37;
Matches 107; Conservative 50; Mismatches 111; Indels 3; Gaps 3;

Qy 4 RSHSLHYLFMGASEQDGLSLFEALGVVDQLFVYDDESRVVEPRTPPWSSRISSQMWL 63
Db 23 RTISLRVFRLOVSDPHGVPEFISGVYDSDHPITTYDSVTRQKBPAPWNAELADHWE 82
Qy 64 QLSQSLKGDHMFVTDFWTIMENHNHKSKE-SHTLQVILGCEMQEDNSTEGYWKYGYDGDH 123
Db 83 RYTQLLRGQWQMFVKELKRLQRHYNHS-GSHTYQRMIGCELLEDGSTTGLQYAYDGDQF 141
Qy 124 LEPCPTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCAQQLLELGRGVLDQ 183
Db 142 LIENKDTLSLWADVNAHTIKQAEANQHLLYQKNWLEBECIAWLKRFLEYGKDTLQRT 201
Qy 184 VPPLVAVKTHVT-SSVTLRCALNYYPQNTIMKWLKQKPMDAKEFEKPDVLPNGDGY 242
Db 202 EPPLVVRNRKETFPFGTALCFKAGFYPPLEYMTWMKNGBEI-VOEIDYDILPVGSGDY 260
Qy 243 QGMITLAVPGEQRYTCQVEHPGLDQPLIV 273
Db 261 QAWASIELDQSSNLXSVCHVEHGVHVLQV 291

RESULT 3

HLRB
MHC class I histocompatibility antigen RLA alpha chain precursor (RL-5) - rabbit
C:Species: Oryctolagus cuniculus (domestic rabbit)
C:Date: 25-Feb-1985 #sequence_revision 25-Feb-1985 #text_change 22-Jun-1999
C:Accession: A02193
R:Tykocinski, M.L.; Marche, P.N.; Max, E.E.; Kindt, T.J.
J. Immunol. 133, 2261-2269, 1984
A:Title: Rabbit class I MHC genes: cDNA clones define full-length transcripts of an expressed gene.
A:Reference number: A02193; MUID:84290724; PMID:6432910
A:Accession: A02193
A:Molecule type: mRNA
A:Residues: 1-361 <TYK>
A:Cross-references: GB:K02441; NID:91293894; PIDN:AAA98729.1; PID:g165496
A:Note: The source of this protein is a T-lymphoid cell line (RL-5), which has been transferred to the many antigens expressed in mouse (K, D, and L) and human (H) cells.
C:Comment: In contrast to the many antigens expressed in mouse (K, D, and L) and human (H) cells, the RLA alpha chain has a unique structure and complexity.
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
C:Keywords: duplication; glycoprotein; heterodimer; transmembrane protein
F:1-24/Domain: signal sequence #status predicted <SIG>
F:25-361/Product: class I histocompatibility antigen RLA alpha chain #status predicted <EXT>
F:25-307/Domain: extracellular #status predicted <EXT>
F:25-114/Domain: alpha-1 <EX1>
F:115-206/Domain: alpha-2 <EX2>
F:220-285/Domain: immunoglobulin homology <IMM>
F:308-329/Domain: transmembrane #status predicted <TM>
F:330-361/Domain: intracellular #status predicted <INT>

F:110/Binding site: carbohydrate (Asn) (covalent) #status predicted
F:125-188,227-283/Disulfide bonds: #status predicted

Query Match 34.4%; Score 523; DB 1; Length 361;
Best Local Similarity 40.1%; Pred. No. 5.3e-36;
Matches 111; Conservative 44; Mismatches 114; Indels 8; Gaps 7;

Qy 5 SHSLHYLFMGASEQDGLSLFEALGVVDQLFVYDDE--SRVVEPRTPWSSRISSQMW 62
Db 26 SHSMRYFTYSVRPGLEPRFIIVGYVDDTQVFRDSDAASPRMEQAPWM-QQVEPEYW 84
Qy 63 LQLSQLKGDHMFVTDFWTIMENHNHKSKE-SHTLQVILGCEMQEDNSTEGYWKYGYD 120
Db 85 DQQTQIAKDTAQTFRVNLNTALRYYNQSAAGSHTFTQMFGEVWADGRFFHGYQYAYDG 144
Qy 121 QHLEPCPTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCAQQLLELGRGVLD 180
Db 145 ADYIALNEDLRSWTAADTAQNTQKWEAAG-EAERHRAYLRECEVWLRRLYLEMGKETL 203
Qy 181 DQOVPLVAVKTHVTSS-VTTLRCALNYYPQNTIMKWLKQKPMDAKEFEKPDVLPNGD 239
Db 204 QRADPPKAHVTHHPASDREATLRCWALGFYPAEISLTWQDGED-QTQDTLVELVTRPGD 262
Qy 240 GTYQGWITLAVPGEQRYTCQVEHPGLDQPLIVWE 276
Db 263 GTFOKWAAVVVPGEQRYTCRVQHEGLPEPLTLTWE 299

RESULT 4

I46858
MHC class I RLA precursor - rabbit
C:Species: Oryctolagus cuniculus (domestic rabbit)
C:Date: 14-Feb-1997 #sequence_revision 14-Feb-1997 #text_change 21-Jan-2000
C:Accession: I46858
R:Marche, P.N.; Tykocinski, M.L.; Max, E.E.; Kindt, T.J.
Immunogenetics 21, 71-82, 1985
A:Title: Structure of a functional rabbit class I MHC gene: Similarity to human class I A;Reference number: I46858; MUID:85103547; PMID:3917574
A:Accession: I46858
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-361 <MAR>
A:Cross-references: GB:K02819; NID:g165497; PIDN:AAA98730.1; PID:g165498
C:Genetics:
A:Introns: 25/1; 115/1; 207/1; 299/1; 337/1; 348/1
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
F:220-285/Domain: immunoglobulin homology <IMM>

Query Match 34.4%; Score 523; DB 2; Length 361;
Best Local Similarity 40.1%; Pred. No. 5.3e-36;
Matches 111; Conservative 44; Mismatches 114; Indels 8; Gaps 7;

Qy 5 SHSLHYLFMGASEQDGLSLFEALGVVDQLFVYDDE--SRVVEPRTPWSSRISSQMW 62
Db 26 SHSMRYFTYSVRPGLEPRFIIVGYVDDTQVFRDSDAASPRMEQAPWM-QQVEPEYW 84
Qy 63 LQLSQLKGDHMFVTDFWTIMENHNHKSKE-SHTLQVILGCEMQEDNSTEGYWKYGYD 120
Db 85 DQQTQIAKDTAQTFRVNLNTALRYYNQSAAGSHTFTQMFGEVWADGRFFHGYQYAYDG 144
Qy 121 QHLEPCPTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCAQQLLELGRGVLD 180
Db 145 ADYIALNEDLRSWTAADTAQNTQKWEAAG-EAERHRAYLRECEVWLRRLYLEMGKETL 203
Qy 181 DQOVPLVAVKTHVTSS-VTTLRCALNYYPQNTIMKWLKQKPMDAKEFEKPDVLPNGD 239
Db 204 QRADPPKAHVTHHPASDREATLRCWALGFYPAEISLTWQDGED-QTQDTLVELVTRPGD 262
Qy 240 GTYQGWITLAVPGEQRYTCQVEHPGLDQPLIVWE 276
Db 263 GTFOKWAAVVVPGEQRYTCRVQHEGLPEPLTLTWE 299

I83063

All.2 - human
 C;Species: Homo sapiens (man)
 C;Date: 02-Aug-1996 #sequence_revision 02-Aug-1996 #text_change 21-Jan-2000
 C;Accession: I83063
 R;Lin, L.; Tokunaga, K.; Ishikawa, Y.; Bannai, M.; Kashiwase, K.; Kuwata, S.; Akaza, T.;
 Tissue Antigens 43, 78-82, 1994
 A;Title: Sequence analysis of serological HLA-A11 split antigens, All.1 and All.2.
 A;Reference number: I60129; MUID:94287401; PMID:8016845

A;Accession: I83063
 A;Status: preliminary; translated from GB/EMBL/DBJ

A;Molecule type: mRNA

A;Residues: 1-365 <RES>

A;Cross-references: GB:D16842; NID:G540517; PIDN:BA04118.1; PID:G487911

C;Genetics:

A;Gene: All02

A;Superfamily: class I histocompatibility antigen; immunoglobulin homology

P;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.9%; Score 515; DB 2; Length 365;
 Best Local Similarity 39.4%; Pred. No. 2.5e-35;
 Matches 109; Conservative 47; Mismatches 113; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDE--SRVPRTPWVSSRISQW 62

DB 26 SHSMRYFYTSVRGRCRPRFIAVGYYDDTQVRFDSDAASQRMPEPRAPWIEQE-GPEYW 84

QY 63 LQLSQSLKGDHMTVDFTWIMENHNHKE-SHTLQVILGCEMQEDNS-TEGWYKYGYDG 120

DB 85 DQETRNVAQSQDTRVDLGLTRGYNQSDGSHTIQIMYCGDVPGRFLGRYQDAYDG 144

QY 121 QDHLFCFPTLDWRAEPRAPWPKLEWERHKIRARONRAYLERDCPAQQLLELGRVYL 180

DB 145 KDYALNEDLRSTWTAADMAAQITKRWAAH-AAEQRAYLEGRCVLEWLRYLENGKETL 203

QY 181 DQVPPPLVKVTHH-VTSSVTLRCALNYYPNITMKWLDKQPMDAKEPEKDVLPNGD 239

DB 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTVELVETRPAGD 262

QY 240 GTYQGWITLAVPPGEORYTCQVHPGLDPLIVWE 276

DB 263 GTFQKAAVVPVSGEORYTCHVQHEGLPKPLTRWE 299

RESULT 9

A47636

MHC class I histocompatibility antigen HLA-A11 alpha chain precursor - human

C;Species: Homo sapiens (man)

C;Date: 31-Dec-1993 #sequence_revision 28-Apr-1995 #text_change 23-Jul-1999

C;Accession: S03536; S03694; A47636; I60129

R;Mayer, W.E.; Jonker, M.; Klein, D.; Ivanyi, P.; van Seventer, G.; Klein, J.

EMBO J. 7, 2765-2774, 1988

A;Title: Nucleotide sequences of chimpanzee MHC class I alleles: evidence for trans-spec

A;Reference number: S01171; MUID:89030641; PMID:2460344

A;Accession: S03536

A;Molecule type: mRNA

A;Residues: 1-365 <MAY>

A;Cross-references: EMBL:X13111; NID:G32138; PIDN:CAA31503.1; PID:G32139

A;Note: this allele is designated A*1101 (formerly AllE, All.1)

A;Accession: S03694

A;Molecule type: mRNA

A;Residues: 1-42, 'K', 44-298 <MA2>

A;Cross-references: EMBL:X13112; NID:G32142; PIDN:CAA31504.1; PID:G32143

A;Note: this allele is designated A*1102 (formerly AllK, All.2)

R;Cowan, E.P.; Jelachich, M.L.; Biddison, W.E.; Colligan, J.E.

Immunogenetics 25, 241-250, 1987

A;Title: DNA sequence of HLA-A11: remarkable homology with HLA-A3 allows identification

A;Reference number: A47636; MUID:87192928; PMID:2437024

A;Accession: A47636

A;Molecule type: DNA

A;Residues: 26-365 <COW>

A;Cross-references: GB:M16007; GB:M16008; GB:M16009; GB:M16010; NID:G184130; PIDN:AAA654

A;Note: the authors translated the codon GAC for residue 89 as Ala, CCG for residue 104

A;Note: this allele is designated A*1101 (formerly AllE, All.1)

R;Lin, L.; Tokunaga, K.; Ishikawa, Y.; Bannai, M.; Kashiwase, K.; Kuwata, S.; Akaza, T.

Tissue Antigens 43, 78-82, 1994

A;Title: Sequence analysis of serological HLA-A11 split antigens, All.1 and All.2.

A;Reference number: I60129; MUID:94287401; PMID:8016845

A;Accession: I60129

A;Status: preliminary; translated from GB/EMBL/DBJ

A;Molecule type: mRNA

A;Residues: 1-365 <RES>

A;Cross-references: GB:D16841; NID:G540516; PIDN:BA04117.1; PID:G487909

A;Note: this allele is designated A*1101 (formerly AllE, All.1)

C;Genetics:

A;Gene: GDB:HLA-A

A;Cross-references: GDB:119310; OMIM:142800

A;Map position: 6p21.3-6p21.3

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

C;Keywords: transmembrane protein

F;1-24/Domain: signal sequence #status predicted <SIG>

F;25-365/Product: class I histocompatibility antigen alpha chain #status predicted <MAT

F;25-298/Domain: extracellular #status predicted <EXT>

F;220-285/Domain: immunoglobulin homology <IMM>

F;239-337/Domain: transmembrane #status predicted <TM>

F;338-365/Domain: intracellular #status predicted <INT>

Query Match 33.8%; Score 514; DB 2; Length 365;

Best Local Similarity 39.4%; Pred. No. 3e-35;

Matches 109; Conservative 47; Mismatches 113; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDE--SRVPRTPWVSSRISQW 62

DB 26 SHSMRYFYTSVRGRCRPRFIAVGYYDDTQVRFDSDAASQRMPEPRAPWIEQE-GPEYW 84

QY 63 LQLSQSLKGDHMTVDFTWIMENHNHKE-SHTLQVILGCEMQEDNS-TEGWYKYGYDG 120

DB 85 DQETRNVAQSQDTRVDLGLTRGYNQSDGSHTIQIMYCGDVPGRFLGRYQDAYDG 144

QY 121 QDHLFCFPTLDWRAEPRAPWPKLEWERHKIRARONRAYLERDCPAQQLLELGRVYL 180

DB 145 KDYALNEDLRSTWTAADMAAQITKRWAAH-AAEQRAYLEGRCVLEWLRYLENGKETL 203

QY 181 DQVPPPLVKVTHH-VTSSVTLRCALNYYPNITMKWLDKQPMDAKEPEKDVLPNGD 239

DB 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTVELVETRPAGD 262

QY 240 GTYQGWITLAVPPGEORYTCQVHPGLDPLIVWE 276

DB 263 GTFQKAAVVPVSGEORYTCHVQHEGLPKPLTRWE 299

RESULT 10

HLA-A30.3 precursor - human

C;Species: Homo sapiens (man)

C;Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 21-Jan-2000

C;Accession: I56039

R;Kato, K.; Trapani, J.A.; Alloppenna, J.; Dupont, B.; Yang, S.Y.

J. Immunol. 143, 3371-3378, 1989

A;Title: Molecular analysis of the serologically defined HLA-Aw19 antigens. A genetical.

A;Reference number: I56039; MUID:90038496; PMID:2478623

A;Accession: I56039

A;Status: preliminary; translated from GB/EMBL/DBJ

A;Molecule type: DNA

A;Residues: 1-365 <RES>

A;Cross-references: GB:M30576; NID:G187646; PIDN:AAA59612.1; PID:G386878

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.8%; Score 514; DB 2; Length 365;

Best Local Similarity 39.4%; Pred. No. 3e-35;

Matches 109; Conservative 48; Mismatches 112; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDE--SRVPRTPWVSSRISQW 62

DB 26 SHSMRYFYTSVRGRCRPRFIAVGYYDDTQVRFDSDAASQRMPEPRAPWIEQE-GPEYW 84

QY 63 LQLSQSLKGDHMTVDFTWIMENHNHKE-SHTLQVILGCEMQEDNS-TEGWYKYGYDG 120

DB 85 DQETRNVAQSQDTRVDLGLTRGYNQSDGSHTIQIMYCGDVPGRFLGRYQDAYDG 144

QY 121 QDHLFCFPTLDWRAEPRAPWPKLEWERHKIRARONRAYLERDCPAQQLLELGRVYL 180

DB 145 KDYALNEDLRSTWTAADMAAQITKRWAAH-AAEQRAYLEGRCVLEWLRYLENGKETL 203

QY 181 DQVPPPLVKVTHH-VTSSVTLRCALNYYPNITMKWLDKQPMDAKEPEKDVLPNGD 239

DB 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTVELVETRPAGD 262

A;Note: submitted to the EMBL Data Library, November 1994

C;Genetics:

A;Gene: hla-A

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology
F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.5%; Score 509; DB 2; Length 365;
Best Local Similarity 39.4%; Pred. No. 7.9e-35;
Matches 109; Conservative 46; Mismatches 114; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDE--SRVPRTPWVSSRISSQMW 62
DB 26 SHSMRYEFTSVSRGPRGFIAVGYVDDTQFVRFSDAASQRMPEAPWIEQE-GPEYW 84
QY 63 LQLSQSLKGDHMFVTDFWTIMENHNHNSKE-SHTLQVILGCEMQED-NSTEGYWKYGYDG 120
DB 85 DGETRKVKAKHSQTHRVDLGLTGLRGYNSQSEAGSHTVQRMVCGDVGSDWRFLRGYHQYAYDG 144
QY 121 QDHLFCFCDPTLDWRAAEPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 145 KDVIYALKEDLRSWTAADMAAQTTHKWEAAHV-AEQRAYLEGCEVWLRRLYLENGKETL 203
QY 181 DQOVPLVKVTHH-VTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEFEKDVLPNGD 239
DB 204 QRTDAPKTHMTHAVSDHEATLRCWALSFPYPAEITLTWQDGED-OTQDTVELVETRPAGD 262
QY 240 GTYQGWITLAVPPEGEORYTCQVEHPGLDQPLIVWE 276
DB 263 GTFOKWAADVVPVSGQEQRYTCHVQHEGLPKPLTLRWE 299

RESULT 14

I38442

gene HLA-A-0205 protein - human

C;Species: Homo sapiens (man)

C;Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 21-Jan-2000

C;Accession: I38442

R;Holmes, N.; Ennis, P.; Wan, A.M.; Denney, D.W.; Parham, P.

J. Immunol. 139, 936-941, 1987

A;Title: Multiple genetic mechanisms have contributed to the generation of the HLA-A2/A24

A;Reference number: I38441; MUID: 87252273; PMID: 3496393

A;Accession: I38442

A;Status: preliminary; translated from GB/EMBL/DBJ

A;Molecule type: DNA

A;Residues: 1-365 <RES>

A;Cross-references: EMBL:U03862; NID:9432436; PIDN:ARA03603.1; PID:9432437

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.5%; Score 509; DB 2; Length 365;
Best Local Similarity 39.7%; Pred. No. 7.9e-35;
Matches 110; Conservative 44; Mismatches 115; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDE--SRVPRTPWVSSRISSQMW 62
DB 26 SHSMRYEFTSVSRGPRGFIAVGYVDDTQFVRFSDAASQRMPEAPWIEQE-GPEYW 84
QY 63 LQLSQSLKGDHMFVTDFWTIMENHNHNSKE-SHTLQVILGCEMQED-NSTEGYWKYGYDG 120
DB 85 DGETRKVKAKHSQTHRVDLGLTGLRGYNSQSEAGSHTVQRMVCGDVGSDWRFLRGYHQYAYDG 144
QY 121 QDHLFCFCDPTLDWRAAEPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 145 KDVIYALKEDLRSWTAADMAAQTTHKWEAAHV-AEQRAYLEGCEVWLRRLYLENGKETL 203
QY 181 DQOVPLVKVTHH-VTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEFEKDVLPNGD 239
DB 204 QRTDAPKTHMTHAVSDHEATLRCWALSFPYPAEITLTWQDGED-OTQDTVELVETRPAGD 262
QY 240 GTYQGWITLAVPPEGEORYTCQVEHPGLDQPLIVWE 276
DB 263 GTFOKWAADVVPVSGQEQRYTCHVQHEGLPKPLTLRWE 299

RESULT 15

I61902

MHC class I histocompatibility antigen HLA-A alpha chain precursor - human (isolate A*0201)

C;Species: Homo sapiens (man)

A;Variety: isolate A*0212

C;Date: 06-Sep-1996 #sequence_revision 06-Sep-1996 #text_change 23-Jul-1999

C;Accession: I61902

R;Belich, M.P.; Madrigal, J.A.; Hildebrand, W.H.; Zemmour, J.; Williams, R.C.; Luz, R.;

Nature 357, 326-329, 1992

A;Title: Unusual HLA-B alleles in two tribes of Brazilian Indians.

A;Reference number: I37120; MUID:92269955; PMID:1317015

A;Accession: I61902

A;Status: translated from GB/EMBL/DBJ

A;Molecule type: mRNA

A;Residues: 1-365 <RES>

A;Cross-references: GB:M84378; NID:9187625; PIDN:AAA59604.1; PID:9187626

A;Experimental source: cell line KRC 033; isolate A*0212.

C;Genetics:

A;Gene: GDB:HLA-A

A;Cross-references: GDB:119310; OMIM:142800

A;Map position: 6p21.3-6p21.3

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

C;Keywords: transmembrane protein

F;1-24/Domain: signal sequence #status predicted <SIG>

F;25-365/Product: MHC class I histocompatibility antigen HLA-A alpha chain #status pred.

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.5%; Score 509; DB 2; Length 365;
Best Local Similarity 39.4%; Pred. No. 7.9e-35;
Matches 109; Conservative 45; Mismatches 115; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDE--SRVPRTPWVSSRISSQMW 62
DB 26 SHSMRYEFTSVSRGPRGFIAVGYVDDTQFVRFSDAASQRMPEAPWIEQE-GPEYW 84
QY 63 LQLSQSLKGDHMFVTDFWTIMENHNHNSKE-SHTLQVILGCEMQED-NSTEGYWKYGYDG 120
DB 85 DGETRKVKAKHSQTHRVDLGLTGLRGYNSQSEAGSHTVQRMVCGDVGSDWRFLRGYHQYAYDG 144
QY 121 QDHLFCFCDPTLDWRAAEPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 145 KDVIYALKEDLRSWTAADMAAQTTHKWEAAHV-AEQRAYLEGCEVWLRRLYLENGKETL 203
QY 181 DQOVPLVKVTHH-VTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEFEKDVLPNGD 239
DB 204 QRTDAPKTHMTHAVSDHEATLRCWALSFPYPAEITLTWQDGED-OTQDTVELVETRPAGD 262
QY 240 GTYQGWITLAVPPEGEORYTCQVEHPGLDQPLIVWE 276
DB 263 GTFOKWAADVVPVSGQEQRYTCHVQHEGLPKPLTLRWE 299

Search completed: August 5, 2003, 13:10:36

Job time : 16 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:05:29 ; Search time 9.5 Seconds
(without alignments)
1366.250 Million cell updates/sec

Title: US-10-092-404-2
Perfect score: 1520
Sequence: 1 RLLRSHLYFLMGASEQDL.....RYTCQVEHGLDQPLIVWE 276

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 127863 seqs, 47026705 residues

Total number of hits satisfying chosen parameters: 127863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : SwissProt_41.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1513	99.5	348	1 HFE_HUMAN	Q30201 homo sapien
2	1238	81.4	348	1 HFE_DICSU	Q9G142 dicorothinu
3	1236	81.3	348	1 HFE_CERSI	Q9GK20 ceratotheri
4	1232	81.1	348	1 HFE_RHUIO	Q9G141 rhinoceros
5	1229	80.9	348	1 HFE_DICBI	Q9G143 diceros bic
6	1156	76.1	360	1 HFE_RAT	Q35799 rattus norv
7	1140	75.0	359	1 HFE_MOUSE	P70387 mus musculu
8	523	34.4	361	1 HALA_RABIT	P01894 oryctolagus
9	523	34.4	361	1 HALB_RABIT	P06140 oryctolagus
10	517	34.0	365	1 LA01_PANTR	P16209 pan troglod
11	516	33.9	364	1 HALB_BOVIN	P13753 bos taurus
12	514	33.8	365	1 LA11_HUMAN	P13746 homo sapien
13	512	33.7	370	1 LA03_HUMAN	P04439 homo sapien
14	510	33.6	365	1 LA80_HUMAN	Q09160 homo sapien
15	508	33.4	365	1 LA02_HUMAN	P16189 homo sapien
16	506	33.3	365	1 LA02_HUMAN	P01892 homo sapien
17	506	33.3	365	1 LA30_HUMAN	P16188 homo sapien
18	506	33.3	365	1 LA74_HUMAN	P30459 homo sapien
19	504	33.2	365	1 LA03_PANTR	P13748 pan troglod
20	503	33.1	365	1 LA33_HUMAN	P16190 homo sapien
21	503	33.1	365	1 LA36_HUMAN	P30455 homo sapien
22	503	33.1	365	1 LA68_HUMAN	P01891 homo sapien
23	501.5	33.0	362	1 HAL9_CANFA	P18466 canis famil
24	501	33.0	365	1 LA01_HUMAN	P30443 homo sapien
25	500	32.9	273	1 LA69_HUMAN	P10316 homo sapien
26	500	32.9	365	1 LA04_PANTR	P13749 pan troglod
27	500	32.9	365	1 LA24_HUMAN	P05534 homo sapien
28	498	32.7	360	1 HALA_BOVIN	P13752 bos taurus
29	497	32.7	296	1 ZA2G_RAT	Q63678 rattus norv
30	497	32.7	362	1 LB45_HUMAN	P30485 homo sapien
31	496	32.6	365	1 LA23_HUMAN	P30447 homo sapien
32	494	32.5	338	1 LB20_HUMAN	P30467 homo sapien
33	493	32.4	363	1 LB04_GORGO	P30382 gorilla gor

34	492	32.4	295	1 ZA2G_HUMAN	P25311 homo sapien
35	492	32.4	322	1 HA10_MOUSE	P01898 mus musculu
36	492	32.4	362	1 LB29_HUMAN	P18463 homo sapien
37	492	32.4	371	1 HA12_RAT	P16391 rattus norv
38	491	32.3	365	1 LA34_HUMAN	P30453 homo sapien
39	491	32.3	365	1 LA66_HUMAN	P30457 homo sapien
40	490	32.2	338	1 HL4G_HUMAN	P17893 homo sapien
41	490	32.2	361	1 LB14_HUMAN	P30389 homo sapien
42	490	32.2	362	1 LB18_HUMAN	P10318 homo sapien
43	490	32.2	366	1 IC02_GORGO	P30385 gorilla gor
44	490	32.2	366	1 IC04_GORGO	P30387 gorilla gor
45	489	32.2	359	1 LB01_PANTR	P13750 pan troglod

ALIGNMENTS

RESULT 1
HFE_HUMAN
ID HFE_HUMAN STANDARD; PRT; 348 AA.
AC Q30201; O75929; O75930; O75931; Q96KU5; Q96KU7; Q96KU8; Q96KU9; Q96KU64;
AC Q9HC68; Q9HC70; Q9HC83;
DT 01-NOV-1997 (Rel. 35, Created)
DT 01-NOV-1997 (Rel. 35, Last sequence update)
DT 15-SEP-2003 (Rel. 42, Last annotation update)
DE Hereditary hemochromatosis protein precursor (HLA-H).
GN HFE OR HLAH.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OC NCBI_TaxId=9606;
RX [1]
RX SEQUENCE FROM N.A. (ISOFORM 1), AND VARIANTS HH ASP-63 AND TYR-282.
RX MEDLINE=96331279; PubMed=8696333;
RA Feder J.N., Gnikre A., Thomas W., Tsuchihashi Z., Ruddy D.A.,
RA Hinton L.M., Jones N.L., Kimmel B.E., Domingo R., Ellis M.C., Jr., Fullan A.,
RA Basava A., Dornishian F., Moore T., Morikang E., Meyer N.C.,
RA Lee V.K., Loeb D.B., Mapa F.A., McClelland E., Fuller A.,
RA Mintier G.A., Moeller N., Moore T., Morikang E., Prass C.E.,
RA Quintana L., Starnes S.M., Schatzman R.C., Brunke K.J.,
RA Drayna D.T., Risch N.J., Bacon B.R., Wolff R.K.;
RA "A novel MHC class I-like gene is mutated in patients with hereditary
haemochromatosis.";
RL Nat. Genet. 13:399-409(1996).
RX SEQUENCE FROM N.A. (ISOFORM 1).
RA Albright W., Burnester N., Bode C., Doenecke D., Drabent B.;
RA Submitted (MAR-1997) to the EMBL/GenBank/DBJ databases.
RX SEQUENCE FROM N.A. (ISOFORM 1).
RX MEDLINE=97294057; PubMed=9149941;
RA Ruddy D.A., Kronmal G.S., Lee V.K., Mintier G.A., Quintana L.,
RA Domingo R., Jr., Meyer N.C., Irrinki A., McClelland E.E., Fullan A.,
RA Mapa F.A., Moore T., Thomas W., Loeb D.B., Harmon C., Tsuchihashi Z.,
RA Wolff R.K., Schatzman R.C., Feder J.N.;
RA "A 1.1-Mb transcript map of the hereditary hemochromatosis locus.";
RL Genome Res. 7:441-456(1997).
RX SEQUENCE FROM N.A. (ISOFORM 1).
RA Gasparini P.;
RA Submitted (SEP-1997) to the EMBL/GenBank/DBJ databases.
RX SEQUENCE FROM N.A. (ISOFORMS 2; 3 AND 4).
RX MEDLINE=99180629; PubMed=10079302;
RA Rhodes D.A., Trowdale J.;
RT "Alternate splice variants of the hemochromatosis gene Hfe.";
RL Immunogenetics 49:357-359(1999).
RX SEQUENCE FROM N.A. (ISOFORMS 2; 5; 6 AND 7).
RA Oliva R., Sanchez M.;
RT "Identification of different alternative splicing forms of the HFE
gene.";
RL Submitted (SEP-2001) to the EMBL/GenBank/DBJ databases.

RN [7] SEQUENCE FROM N.A. (ISOFORMS 1; 7; 8; 9 AND 10).
 RP MEDLINE=20448010; PubMed=11001625;
 RA Thénie A., Orhant M., Gicquel I., Fergelot P., Le Gall J.-Y.,
 RA David V., Mosser J.;
 RT "The HFE gene undergoes alternate splicing processes.";
 RL Blood Cells Mol. Dis. 26:155-162(2000).
 RN [8]
 RN FUNCTION.
 RP MEDLINE=98132614; PubMed=9465039;
 RA Feder J.N., Penny D.M., Irfink A., Lee V.K., Lebron J.A., Watson N.,
 RA Tsuchihashi Z., Sigal E., Bjorkman P.J., Schatzman R.C.;
 RT "The hemochromatosis gene product complexes with the transferrin
 RT receptor and lowers its affinity for ligand binding.";
 RL Proc. Natl. Acad. Sci. U.S.A. 95:1472-1477(1998).
 RN [9]
 RP X-RAY CRYSTALLOGRAPHY (2.6 ANGSTROMS).
 RP MEDLINE=98206473; PubMed=9546397;
 RA Lebron J.A., Bennett M.J., Vaughn D.E., Chirino A.J., Snow P.M.,
 RA Mintier G.A., Feder J.N., Bjorkman P.J.;
 RT "Crystal structure of the hemochromatosis protein HFE and
 RT characterization of its interaction with transferrin receptor.";
 RL Cell 93:111-123(1998).
 RN [10]
 RP VARIANTS HH ASP-63 AND TYR-282.
 RP MEDLINE=97260408; PubMed=9106528;
 RA Carella M., D'Ambrosio L., Totaro A., Grifa A., Valentino M.A.,
 RA Piperno A., Girelli D., Roetto A., Franco B., Gasparini P.,
 RA Canaschella C.;
 RT "Mutation analysis of the HLA-H gene in Italian hemochromatosis
 RT patients.";
 RL Am. J. Hum. Genet. 60:828-832(1997).
 RN [11]
 RP VARIANT HH/PCT TYR-282.
 RP MEDLINE=97176837; PubMed=9024376;
 RA Roberts A.G., Whitley S.D., Morgan R.R., Worwood M., Elder G.H.;
 RT "Increased frequency of the hemochromatosis Cys282Tyr mutation in
 RT sporadic porphyria cutanea tarda.";
 RL Lancet 349:321-323(1997).
 RN [12]
 RP VARIANTS HH/PCT ASP-63.
 RP MEDLINE=98085904; PubMed=9425935;
 RA Sampletro M., Piperno A., Lupica L., Arosio C., Vergani A.,
 RA Corbetta N., Malosio I., Mattioli M., Fracanzani A.L.,
 RA Cappellini M.D., Fiorelli G., Fargion S.;
 RT "High prevalence of the His63Asp HFE mutation in Italian patients with
 RT porphyria cutanea tarda.";
 RL Hepatology 27:181-184(1998).
 RN [13]
 RP VARIANTS HH/PCT ASP-63 AND TYR-282.
 RP MEDLINE=98281650; PubMed=9620340;
 RA Bonkovsky H.L., Poh-Fitzpatrick M., Pinestone N., Obando J.,
 RA Di Biasele A., Tattire C., Tortorelli K., LeClair P., Mercurio M.G.,
 RA Lambrecht R.W.;
 RT "Porphyria cutanea tarda, hepatitis C, and HFE gene mutations in North
 RT America";
 RL Hepatology 27:1661-1669(1998).
 RN [14]
 RP VARIANTS HH ASP-63; CYS-65 AND TYR-282.
 RP MEDLINE=99211934; PubMed=10194428;
 RA Mura C., Raguene O., Ferec C.;
 RT "HFE mutations analysis in 711 hemochromatosis probands: evidence for
 RT S65C implication in mild form of hemochromatosis.";
 RL Blood 93:2502-2505(1999).
 RN [15]
 RP VARIANTS HH CYS-65; ARG-93 AND THR-105.
 RP MEDLINE=20042794; PubMed=10575540;
 RA Barton J.C., Sawada-Hirai R., Rothenberg B.E., Acton R.T.;
 RT "Two novel missense mutations of the HFE gene (I105T and G93R) and
 RT identification of the S65C mutation in Alabama hemochromatosis
 RT probands";
 RL Blood Cells Mol. Dis. 25:147-155(1999).
 RN [16]

RP VARIANTS VP ASP-63 AND HIS-127, VARIANT HH MET-330, AND VARIANTS
 RP MET-53 AND MET-59.
 RX MEDLINE=99330560; PubMed=10401000;
 RA de Villiers J.N.P., Hillermann R., Loubser L., Kotze M.J.;
 RT "Spectrum of mutations in the HFE gene implicated in haemochromatosis
 RT and porphyria";
 RL Hum. Mol. Genet. 8:1517-1522(1999).
 RN [17]
 RP VARIANTS HH ASP-63 AND TYR-282.
 RP MEDLINE=99140360; PubMed=10094552;
 RA Merryweather-Clarke A.T., Simonsen H., Shearman J.D., Poynton J.J.,
 RA Norgaard-Pedersen B., Robson K.J.H.;
 RT "A retrospective anonymous pilot study in screening newborns for HFE
 RT mutations in Scandinavian populations.";
 RL Hum. Mutat. 13:154-159(1999).
 RN [18]
 RP VARIANT HH CYS-65.
 RA Fagan E., Payne S.J.;
 RT "A novel missense mutation S65C in the HFE gene with a possible role
 RT in hereditary haemochromatosis.";
 RL Hum. Mutat. 13:507-508(1999).
 RN [19]
 RP VARIANT LYS-277.
 RP MEDLINE=20081073; PubMed=10612845;
 RA Bradbury R., Fagan E., Payne S.J.;
 RT "Two novel polymorphisms (E277K and V212V) in the haemochromatosis
 RT gene HFE.";
 RL Hum. Mutat. 15:120-120(2000).
 CC -1- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
 CC affinity for iron-loaded transferrin.
 CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -1- ALTERNATIVE PRODUCTS:
 CC Event=Alternative splicing; Named isoforms=10;
 CC Comment=Additional isoforms seem to exist;
 CC
 CC Name=1;
 CC IsoId=Q30201-1; Sequence=Displayed;
 CC Name=2; Synonyms=delE2;
 CC IsoId=Q30201-2; Sequence=VSP_003218;
 CC Name=3; Synonyms=del14E4;
 CC IsoId=Q30201-3; Sequence=VSP_003225;
 CC Name=4; Synonyms=delE214E4;
 CC IsoId=Q30201-4; Sequence=VSP_003218, VSP_003225;
 CC Name=5;
 CC IsoId=Q30201-5; Sequence=VSP_003219;
 CC Name=6;
 CC IsoId=Q30201-6; Sequence=VSP_003220;
 CC Name=7; Synonyms=delE3;
 CC IsoId=Q30201-7; Sequence=VSP_003221;
 CC Name=8; Synonyms=1043-2283del,intron6ins;
 CC IsoId=Q30201-8; Sequence=VSP_003226, VSP_003227;
 CC Name=9; Synonyms=delE3-7;
 CC IsoId=Q30201-9; Sequence=VSP_003223, VSP_003224;
 CC Name=10; Synonyms=582-878del;
 CC IsoId=Q30201-10; Sequence=VSP_003222;
 CC -1- TISSUE SPECIFICITY: IN ALL TISSUES TESTED EXCEPT BRAIN.
 CC -1- DISEASE: DEFECTS IN HFE ARE A CAUSE OF HEREDITARY HEMOCHROMATOSIS
 CC (HH). HH IS AN AUTOSOMAL RECESSIVE INBORN DISORDER OF IRON
 CC METABOLISM. FREQUENT AMONG CAUCASIANS. HH IS CHARACTERIZED BY
 CC ABNORMAL INTESTINAL IRON ABSORPTION AND PROGRESSIVE INCREASE OF
 CC TOTAL BODY IRON, WHICH RESULTS IN MIDLIFE IN CLINICAL
 CC COMPLICATIONS INCLUDING CIRRHOSIS, CARDIOPATHY, DIABETES,
 CC ENDOCRINE DYSFUNCTIONS, ARTHROPATHY, AND SUSCEPTIBILITY TO LIVER
 CC CANCER. SINCE THE DISEASE COMPLICATIONS CAN BE EFFECTIVELY
 CC PREVENTED BY REGULAR PHLEBOTOMIES, EARLY DIAGNOSIS IS MOST
 CC IMPORTANT TO PROVIDE A NORMAL LIFE EXPECTANCY TO THE AFFECTED
 CC SUBJECTS.
 CC -1- DISEASE: DEFECTS IN HFE ARE A CAUSE OF PORPHYRIA CUTANEA TARDA
 CC (PCT), A DISORDER CHARACTERIZED BY LIGHT-SENSITIVE DERMATITIS AND
 CC PRESENCE OF LARGE AMOUNTS OF UROPORPHYRIN IN URINE. IRON OVERLOAD
 CC IS OFTEN PRESENT IN ASSOCIATION WITH VARYING DEGREES OF LIVER
 CC DAMAGE. PCT IS THE MOST COMMON FORM OF PORPHYRIA WORLDWIDE. IT
 CC OCCURS IN TWO FORMS: THE SPORADIC TYPE (PCT TYPE I) AND THE
 CC FAMILIAL TYPE (PCT TYPE II), WHICH IS INHERITED IN AN AUTOSOMAL

Query Match 99.5%; Score 1513; DB 1; Length 348;
 Best Local Similarity 99.6%; Pred. No. 1.3e-119;
 Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQDLFFVYDDSRVPRTPWSSRISQ 60
 DB 23 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQDLFFVYDDSRVPRTPWSSRISQ 82

QY 61 MWLQSLKGDHMTFTVDFWTTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
 DB 83 MWLQSLKGDHMTFTVDFWTTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142

QY 121 ODHLEFCPTLDWRAAPRAAPPTKLEWERHKKIRARONRAYLERDCPAQLQQLLELGRGVL 180
 DB 143 ODHLEFCPTLDWRAAPRAAPPTKLEWERHKKIRARONRAYLERDCPAQLQQLLELGRGVL 202

QY 181 DQVPPPLVKVTHVTSVTLRCRALNYYPQNTMKWLKDKQPMDAKEFEPEKDVLPNGDG 240
 DB 203 DQVPPPLVKVTHVTSVTLRCRALNYYPQNTMKWLKDKQPMDAKEFEPEKDVLPNGDG 262

QY 241 TYQGMITLAVPPGEGEORYTCQVEHPGLDQPLIVWE 276
 DB 263 TYQGMITLAVPPGEGEORYTCQVEHPGLDQPLIVWE 298

RESULT 2

HFE_DICSU STANDARD; PRT; 348 AA.

AC Q9GL42;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Hereditary hemochromatosis protein precursor.
 GN HFE.

OS Dicerorhinus sumatrensis (Sumatran rhinoceros).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Perissodactyla; Rhinocerotidae; Dicerorhinus.
 OX NCBI_TaxID=89632;
 [1]
 RN SEQUENCE FROM N.A.
 RA West C.J., Worley M., Beutler E.;
 RT "Rhinoceros HFE polymorphisms."
 RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
 affinity for iron-loaded transferrin.
 CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -1- SIMILARITY: TO MHC CLASS I ANTIGENS.

 CC This SWISS-PROT entry is copyright. It is produced through a collaboration
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 the European Bioinformatics Institute. There are no restrictions on its
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 modified and this statement is not removed. Usage by and for commercial
 entities requires a license agreement (See <http://www.isb-sib.ch/announce/>
 or send an email to license@isb-sib.ch).
 CC EMBL; AY007543; AAG23703.1; --
 CC HSSP; Q30201; 1A6Z.
 DR InterPro; IPR007110; Ig-like.
 DR InterPro; IPR003597; Ig-cl.
 DR InterPro; IPR003006; Ig_MHC.
 DR InterPro; IPR001039; MHC_I.
 DR Pfam; PF00047; Ig_1.
 DR Pfam; PF00129; MHC_I; 1.
 DR ProDom; PD000050; MHC_I; 1.
 DR SMART; SM00407; IGC1; 1.
 DR PROSITE; PS00835; IG_LIKE; 1.
 DR PROSITE; PS00290; IG_MHC; 1.
 DR MHC_I; Transmembrane; Glycoprotein; Signal.
 KW MHC_I; 22 BY SIMILARITY.
 FT SIGNAL 1 22 HEREDITARY HEMOCHROMATOSIS PROTEIN.
 FT CHAIN 23 348
 FT DOMAIN 23 114 EXTRACELLULAR ALPHA-1.

FT DOMAIN 115 205 EXTRACELLULAR ALPHA-2.
 FT DOMAIN 206 297 EXTRACELLULAR ALPHA-3.
 FT DOMAIN 298 306 CONNECTING PEPTIDE.
 FT TRANSMEM 307 330 POTENTIAL.
 FT DOMAIN 331 348 CYTOPLASMIC TAIL.
 FT DISULFID 124 187 BY SIMILARITY.
 FT DISULFID 225 282 BY SIMILARITY.
 FT CARBOHYD 110 110 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 130 130 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 234 234 N-LINKED (GLCNAC. . .) (POTENTIAL).
 SQ SEQUENCE 348 AA; 39740 MW; 518BFD357AB83B90 CRC64;

Query Match 81.4%; Score 1238; DB 1; Length 348;
 Best Local Similarity 81.3%; Pred. No. 1.5e-96;
 Matches 222; Conservative 20; Mismatches 31; Indels 0; Gaps 0;

QY 4 RSHSLHYLFMGASQDGLSLFEALGYVDQDLFFVYDDSRVPRTPWSSRISQMWL 63
 DB 26 RSHSLHYLFMGASQDGLSLFEALGYVDQDLFFVYDDSRVPRTPWSSRISQMWL 85

QY 64 QLSQLKGDHMTFTVDFWTTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDGQDH 123
 DB 86 QLSQLKGDHMTFTVDFWTTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDGQDH 145

QY 124 LEFCPTLDWRAAPRAAPPTKLEWERHKKIRARONRAYLERDCPAQLQQLLELGRGVLDOQ 183
 DB 146 LEFCPTLDWRAAPRAAPPTKLEWERHKKIRARONRAYLERDCPAQLQQLLELGRGVLDOQ 205

QY 184 VPPLVKVTHVTSVTLRCRALNYYPQNTMKWLKDKQPMDAKEFEPEKDVLPNGDGTQY 243
 DB 206 VPPLVKVTHVTSVTLRCRALNYYPQNTMKWLKDKQPMDAKEFEPEKDVLPNGDGTQY 265

QY 244 GWITLAVPPGEGEORYTCQVEHPGLDQPLIVWE 276
 DB 266 SWALAVPPGEGEORYTCQVEHPGLDQPLIVWE 298

RESULT 3

HFE_CERSI STANDARD; PRT; 348 AA.

AC Q9GKZ0;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Hereditary hemochromatosis protein precursor.
 GN HFE.

OS Ceratotherium simum (White rhinoceros) (Square-lipped rhinoceros).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Perissodactyla; Rhinocerotidae; Ceratotherium.
 OX NCBI_TaxID=9807;
 [1]
 RN SEQUENCE FROM N.A.
 RA West C.J., Worley M., Beutler E.;
 RT "Rhinoceros HFE polymorphisms."
 RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
 affinity for iron-loaded transferrin.
 CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -1- SIMILARITY: TO MHC CLASS I ANTIGENS.

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 or send an email to license@isb-sib.ch).
 CC EMBL; AY007541; AAG23701.1; --
 CC HSSP; Q30201; 1A6Z.
 DR InterPro; IPR007110; Ig-like.
 DR InterPro; IPR003597; Ig-cl.
 DR InterPro; IPR003006; Ig_MHC.
 DR InterPro; IPR001039; MHC_I.
 DR Pfam; PF00047; Ig_1.
 DR Pfam; PF00129; MHC_I; 1.
 DR ProDom; PD000050; MHC_I; 1.
 DR SMART; SM00407; IGC1; 1.
 DR PROSITE; PS00835; IG_LIKE; 1.
 DR PROSITE; PS00290; IG_MHC; 1.
 DR MHC_I; Transmembrane; Glycoprotein; Signal.
 KW MHC_I; 22 BY SIMILARITY.
 FT SIGNAL 1 22 HEREDITARY HEMOCHROMATOSIS PROTEIN.
 FT CHAIN 23 348
 FT DOMAIN 23 114 EXTRACELLULAR ALPHA-1.

DR InterPro; IPR001039; MHC_I.
 DR Pfam; PF00047; ig; 1.
 DR Pfam; PF00129; MHC_I; 1.
 DR PRINTS; PR01638; MHCCLASSI.
 DR PRODOM; PD000050; MHC_I; 1.
 DR SMART; SM00407; IGC1; 1.
 DR PROSITE; PS00835; IG_LIKE; 1.
 DR PROSITE; PS00290; IG_MHC; 1.
 KW MHC I; Transmembrane; Glycoprotein; Signal.
 FT SIGNAL 1 22
 FT CHAIN 23 348
 FT DOMAIN 23 114
 FT DOMAIN 115 205
 FT DOMAIN 206 297
 FT DOMAIN 298 306
 FT TRANSMEM 307 330
 FT DOMAIN 331 348
 FT DISULFID 124 187
 FT DISULFID 225 282
 FT CARBOHYD 110 110
 FT CARBOHYD 130 130
 FT CARBOHYD 234 234
 SQ SEQUENCE 348 AA; 39822 MW; 2523016EC9FB9E91 CRC64;
 Query Match 81.3%; Score 1236; DB 1; Length 348;
 Best Local Similarity 81.7%; Pred. No. 2.1e-96;
 Matches 223; Conservative 18; Mismatches 32; Indels 0; Gaps 0;
 QY 4 RSHSLHYLFMGASEQDLGLSLFALGYVDQDLFVYDDSRVRRVPTPWSSRISSQMWL 63
 DB 26 RSHSLRYLFMGASERDGLPLFALGYVDDELFAVYNHESRAESRAQWVLGEAHSQWL 85
 QY 64 QLSQSLKGDWDMFTVDFTIMENHNHSHKESHTLQVILGCEWQEDNSTEGYKYGVDGDH 123
 DB 86 QLSQSLKGDWDMFTVDFTIMDNHNHSHKESHTLQVILGCEVQEDNSTRGFWKYGVDGDH 145
 QY 124 LEFCPDTLDWRAAPRAWPTKLEWRHKIRARQNRAYLERDCPAQLQLLELGRGVLDQ 183
 DB 146 LEFCPETLDWRAAESRALTTKLEWVKNIRAKQNRAYLERDCPEQLQWLLELGRGVLDQ 205
 QY 184 VPPLVKVTHVTSVTLRCRALNYYPQNTWKWKDKQPMDAKEPEKDVLPNGDGTQ 243
 DB 206 VPPLVKVTHVASVTLRCQALNFYQNTWRLKDKRPVMDKDAESKDVLPSPGDGTQ 265
 QY 244 GWITLAVPPGEQRYTCQVEHGLDPLIVIE 276
 DB 266 SWEALAVPPGEQRYTCQVEHGLDPLTATWE 298

RESULT 4
 ID_HFE_RHUN STANDARD; PRIT; 348 AA.
 AC Q9GL41;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Hereditary hemochromatosis protein precursor.
 GN HFE.
 OS Rhinoceros unicornis (Greater Indian rhinoceros).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Perissodactyla; Rhinocerotidae; Rhinoceros.
 OX NCBI_TaxID=9809;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA West C.J., Worley M., Beutler E.;
 RT "Rhinoceros HFE polymorphisms."
 RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
 CC -|- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
 CC affinity for iron-loaded transferrin.
 CC -|- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -|- SIMILARITY: TO MHC CLASS I ANTIGENS.
 CC -----
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 CC -----
 CC EMBL; AY007544; AAG23704.1; -
 DR HSP; Q30201; IASZ.
 DR InterPro; IPR007110; Ig-like.
 DR InterPro; IPR003597; Ig_c1.
 DR InterPro; IPR003006; IG_MHC.
 DR InterPro; IPR001039; MHC_I.
 DR Pfam; PF00047; ig; 1.
 DR Pfam; PF00129; MHC_I; 1.
 DR PRINTS; PR01638; MHCCLASSI.
 DR PRODOM; PD000050; MHC_I; 1.
 DR SMART; SM00407; IGC1; 1.
 DR PROSITE; PS00835; IG_LIKE; 1.
 DR PROSITE; PS00290; IG_MHC; 1.
 KW MHC I; Transmembrane; Glycoprotein; Signal.
 FT SIGNAL 1 22
 FT CHAIN 23 348
 FT DOMAIN 23 114
 FT DOMAIN 115 205
 FT DOMAIN 206 297
 FT DOMAIN 298 306
 FT TRANSMEM 307 330
 FT DOMAIN 331 348
 FT DISULFID 124 187
 FT DISULFID 225 282
 FT CARBOHYD 110 110
 FT CARBOHYD 130 130
 FT CARBOHYD 234 234
 SQ SEQUENCE 348 AA; 39743 MW; F2723D57A327A6B4 CRC64;
 Query Match 81.1%; Score 1232; DB 1; Length 348;
 Best Local Similarity 81.0%; Pred. No. 4.6e-96;
 Matches 221; Conservative 20; Mismatches 32; Indels 0; Gaps 0;
 QY 4 RSHSLHYLFMGASEQDLGLSLFALGYVDQDLFVYDDSRVRRVPTPWSSRISSQMWL 63
 DB 26 RSHSLRYLFMGASERDGLPLFALGYVDDELFAVYNHESRAESRAQWVLGEAHSQWL 85
 QY 64 QLSQSLKGDWDMFTVDFTIMENHNHSHKESHTLQVILGCEWQEDNSTEGYKYGVDGDH 123
 DB 86 QLSQSLKGDWDMFTVDFTIMDNHNHSHKESHTLQVILGCEVQEDNSTRGFWKYGVDGDH 145
 QY 124 LEFCPDTLDWRAAPRAWPTKLEWRHKIRARQNRAYLERDCPAQLQLLELGRGVLDQ 183
 DB 146 LEFCPETLDWRAAESRALTTKLEWVKNIRAKQNRAYLERDCPEQLQWLLELGRGVLDQ 205
 QY 184 VPPLVKVTHVTSVTLRCRALNYYPQNTWKWKDKQPMDAKEPEKDVLPNGDGTQ 243
 DB 206 VPPLVKVTHVASVTLRCQALNFYQNTWRLKDKRPVMDKDAESKDVLPSPGDGTQ 265
 QY 244 GWITLAVPPGEQRYTCQVEHGLDPLIVIE 276
 DB 266 SWALAVPPGEQRYTCQVEHGLDPLTATWE 298
 RESULT 5
 ID_HFE_DICBI STANDARD; PRIT; 348 AA.
 AC Q9GL43; Q9GK81;
 DT 28-FEB-2003 (Rel. 41, Created)
 DT 28-FEB-2003 (Rel. 41, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Hereditary hemochromatosis protein precursor.
 GN HFE.
 OS Dicerus bicornis (Black rhinoceros).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Perissodactyla; Rhinocerotidae; Diceros.


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QY 5 SHSLHYLFMGASEODLGLSFEALGYDDQLFVYDDERRRPRTPWSSRISSQWLQ 64
DB 32 SHSLHYLFMGASEODLGLSFEALGYDDQLFVYDDERRRPRTPWSSRISSQWLQ 91
QY 65 LSQSLKGDHMTVDFTWIMENHNSK-----ESHTLOVILGCEMQEDNSTEGWKY 116
DB 92 LSQSLKGDHMTVDFTWIMENHNSK-----ESHTLOVILGCEMQEDNSTEGWKY 151
QY 117 GYGQDHLFCPTDLWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLEL 176
DB 152 GYGQDHLFCPTDLWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLEL 211
QY 177 RGVLDDQVPLVKVTHVTSVTLRCALNYYPQNTIMKWKDKQPMDAKEPEPKDVL 236
DB 212 RGVLDDQVPLVKVTHVTSVTLRCALNYYPQNTIMKWKDKQPMDAKEPEPKDVL 271
QY 237 NGDGTQGWITLAVPGEORYTCQVHPGLDQPLVIWE 276
DB 272 NGDGTQGWITLAVPGEORYTCQVHPGLDQPLVIWE 311

RESULT 7
HFE_MOUSE
ID HFE_MOUSE STANDARD; PRT; 359 AA.
AC P70387;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Hereditary hemochromatosis protein homolog precursor.
GN HFE OR MR2.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=129/SvJ;
RX MEDLINE=98060831; PubMed=9396865;
RA Riegert P., Gillilan S., Nanda I., Schmid M., Bahram S.;
RT "The mouse HFE gene.";
RL Immunogenetics 47:174-177 (1998).
RN [2]
RP SEQUENCE FROM N.A.
RC STRAIN=BALB/c; TISSUE=Lung;
RA Hashimoto K.;
RL Submitted (SEP-1996) to the EMBL/GenBank/DBJ databases.
RN [3]
RP SEQUENCE OF 37-211 FROM N.A.
RC STRAIN=BALB/c; TISSUE=Liver;
RX MEDLINE=97148566; PubMed=9020055;
RA Hashimoto K., Hirai M., Kurosawa Y.;
RT "Identification of a mouse homolog for the human hereditary
hemochromatosis candidate gene.";
RL Biochem. Biophys. Res. Commun. 230:35-39 (1997).
RN [4]
RP SEQUENCE OF 79-359 FROM N.A.
RC STRAIN=129;
RA Albright W., Drabant B., Donecke D.;
RL Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
affinity for iron-loaded transferrin (By similarity).
CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
CC -----
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CC -----
EMBL; AF007558; AAC03447.1; --

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DR EMBL; U66849; AAB07525.1; --
DR EMBL; Y12650; CAA73197.1; --
DR EMBL; U80604; AAB51504.1; --
DR PIR; JCS382; JCS382.
DR HSSP; Q30201; IA62.
DR MGD; MGI:109191; Hfe.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig_c1.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGC1; 1.
DR PROSITE; PS50835; IG-LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal.
FT SIGNAL 1 24
FT CHAIN 25 359
FT HEREDITARY HEMOCHROMATOSIS PROTEIN
FT HOMOLOG.
FT DOMAIN 25 126
FT DOMAIN 127 217
FT DOMAIN 218 309
FT DOMAIN 310 318
FT TRANSMEM 319 339
FT TRANSMEM 340 359
FT DISULFID 136 139
FT DISULFID 237 294
FT CARBOHYD 114 142
FT CARBOHYD 166 166
FT CARBOHYD 246 246
FT SEQUENCE 359 AA; 40548 MW; 4BDE6C27F9FF20B4 CRC64;
QY 4 RSHSLHYLFMGASEODLGLSFEALGYDDQLFVYDDERRRPRTPWSSRISSQWLQ 63
DB 30 RSHSLHYLFMGASEODLGLSFEALGYDDQLFVYDDERRRPRTPWSSRISSQWLQ 89
QY 64 QLSQSLKGDHMTVDFTWIMENHNSK-----ESHTLOVILGCEMQEDNSTEGWKY 115
DB 90 HLSQSLKGDHMTVDFTWIMENHNSK-----ESHTLOVILGCEMQEDNSTEGWKY 149
QY 116 YGYDQDHLFCPTDLWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLEL 175
DB 150 YGYDQDHLFCPTDLWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLEL 209
QY 176 GRGVLDDQVPLVKVTHVTSVTLRCALNYYPQNTIMKWKDKQPMDAKEPEPKDVL 235
DB 210 GRGVLDDQVPLVKVTHVTSVTLRCALNYYPQNTIMKWKDKQPMDAKEPEPKDVL 269
QY 236 PNGDGTQGWITLAVPGEORYTCQVHPGLDQPLVIWE 276
DB 270 PNGDGTQGWITLAVPGEORYTCQVHPGLDQPLVIWE 310

RESULT 8
HFE_MOUSE
ID HFE_MOUSE STANDARD; PRT; 361 AA.
AC P01894;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE RLA class I histocompatibility antigen, alpha chain 11/11 precursor.
OS Oryctolagus cuniculus (Rabbit).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
OX NCBI_TaxID=9986;
RN [1]

```


QY 121 QDHLFCPTDLDWRAAPRAWPYKLEWERHKIRARONRAYLERDCPAQLOQLLELGRGVL 180
Db 145 ADYIALNEDLRSWTAADTAQNTQKWEAAG-EAERHRAYLRECVLEWRLRYLEMCKETL 203
QY 181 DQVPLVAVTHVTS--VTTLCRALNYPQNTMKWLKDKQPMDAKEEPKDVLPNGD 239
Db 204 QADPPKAVTHHPASDREATLRCWALGFYPAEISLTWQDGED-QTQDTLVELTRPGD 262
QY 240 GTYQGWITLAVPGEQRYTCQVEHPGLDQPLVIVE 276
Db 263 GTQKWAUVVPSGEEQRYTCVQHEGLPEPLTLWE 299
RESULT 10
1A01 PANTR
ID 1A01 PANTR STANDARD; PRT; 365 AA.
AC P16209;
DT 01-APR-1990 (Rel. 14, Created)
DT 01-APR-1990 (Rel. 14, Last sequence update)
DT 01-APR-1993 (Rel. 25, Last annotation update)
DE CHLA class I histocompatibility antigen, A-2 alpha chain precursor.
OS Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
OX NCBI_TaxID=9598;
RN [1]
SEQUENCE FROM N.A.
RX MEDLINE=90201944; PubMed=1690682;
RA Lawlor D.A., Warren E., Ward P.E., Parham P.;
RT "Comparison of class I MHC alleles in humans and apes";
RL Immunol. Rev. 113:147-185 (1990).
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
THE IMMUNE SYSTEM.
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
MICROGLOBULIN).
CC
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CC
CC EMBL; M30678; AAA87970.1; --
DR PIR; I36961; I36961.
DR HSP; Q95352; IHK.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig_cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR PRODOM; PD000050; MHC_I; 1.
DR SMART; SM00407; IGC1; 1.
DR PROSITE; PS50835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal.
FT SIGNAL 1 24
FT CHAIN 25 365 CHLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
A-2 ALPHA CHAIN.
FT FT
FT DOMAIN 25 114 EXTRACELLULAR ALPHA-1.
FT DOMAIN 115 206 EXTRACELLULAR ALPHA-2.
FT DOMAIN 207 298 EXTRACELLULAR ALPHA-3.
FT DOMAIN 299 308 CONNECTING PEPTIDE.
FT TRANSMEM 309 332
FT DOMAIN 333 365 CYTOPLASMIC TAIL.
FT DISULFID 125 188 BY SIMILARITY.
FT DISULFID 227 283 BY SIMILARITY.
FT CARBOHYD 110 110 N-LINKED (GLNAC...) (BY SIMILARITY).
SQ SEQUENCE 365 AA; 40848 MW; FC45276BD038D3E CRC64;

Query Match 34.0%; Score 517; DB 1; Length 365;
Best Local Similarity 39.7%; Pred. No. 3.9e-36;
Matches 110; Conservative 45; Mismatches 114; Indels 8; Gaps 7;
QY 5 SLSHLVFMGASEQDGLSLFEALGVVDOLFVYDDE--SRVERPTWVSSRISSOMW 62
Db 26 SHSMRYFFTSVSRPGGEPRFIAVGVDVDTQVFRFSDAASQRMPEAPRAWIEQE-GPEYW 84
QY 63 LQLSQLKGDHMFVDFWTIMENHNHNSKE-SHTLQVILGCEMQEDNS-TEGWYKTYDYG 120
Db 85 DEETRSKAKAHSQTDRLDGLTGRGYNQSEDSHTIQTIMYGCDSGRFLRGVRYDAYDG 144
QY 121 QDHLFCPTDLDWRAAPRAWPYKLEWERHKIRARONRAYLERDCPAQLOQLLELGRGVL 180
Db 145 KDYIALNEDLRSWTAADTAQNTQKWEAAG-EAERHRAYLRECVLEWRLRYLEMCKETL 203
QY 181 DQVPLVAVTHVTS--VTTLCRALNYPQNTMKWLKDKQPMDAKEEPKDVLPNGD 239
Db 204 QADPPKAVTHHPASDREATLRCWALGFYPAEISLTWQDGED-QTQDTLVELTRPGD 262
QY 240 GTYQGWITLAVPGEQRYTCQVEHPGLDQPLVIVE 276
Db 263 GTQKWAUVVPSGEEQRYTCVQHEGLPEPLTLWE 299
RESULT 11
HA1B BOVIN STANDARD; PRT; 364 AA.
ID HA1B BOVIN STANDARD; PRT; 364 AA.
AC P13753;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE BOLA class I histocompatibility antigen, alpha chain BL3-7 precursor.
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxID=9913;
RN [1]
SEQUENCE FROM N.A.
RX MEDLINE=88258075; PubMed=3133413;
RA Emis P.D., Jackson A.P., Parham P.;
RT "Molecular cloning of bovine class I MHC cDNA";
RL J. Immunol. 141:642-651 (1988).
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
THE IMMUNE SYSTEM.
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
MICROGLOBULIN).
CC
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CC
CC EMBL; M21043; AAA30641.1; --
DR HSP; P16391; LED3.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig_cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR PRODOM; PD000050; MHC_I; 1.
DR SMART; SM00407; IGC1; 1.
DR PROSITE; PS50835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal.
FT SIGNAL 1 27
FT CHAIN 28 364 BOLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT FT

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FT DOMAIN 28 117 ALPHA CHAIN B13-7.
FT DOMAIN 118 209 EXTRACELLULAR ALPHA-1.
FT DOMAIN 210 301 EXTRACELLULAR ALPHA-2.
FT DOMAIN 302 310 EXTRACELLULAR ALPHA-3.
FT TRANSMEM 311 331 CONNECTING PEPTIDE.
FT DOMAIN 332 364 CYTOPLASMIC.
FT CARBOHYD 106 106 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 113 113 N-LINKED (GLCNAC. . .) (BY SIMILARITY).
FT DISULFID 128 191 BY SIMILARITY.
FT DISULFID 230 286 BY SIMILARITY.
SQ SEQUENCE 364 AA; 41513 MW; 622056CF7DCE7873 CRC64;

Query Match 33.9%; Score 516; DB 1; Length 364;
Best Local Similarity 38.9%; Pred. No. 4.8e-36;
Matches 109; Conservative 50; Mismatches 113; Indels 8; Gaps 7;

QY 2 LLRSLSLHLYFMGASEQDGLSLFEALGVDDQLFVYDDB--SRVRPRTWVSSRISS 59
DB 26 LAGSHSLURYPTGTGVSRRGLGEPRPIAVGVYDDTQFVFRSDAPNPREPVPWMEQSGP 84
QY 60 QMWLQSLQSLKGDHMTVDFTWIMENHNHKSKE-SHTLQVILGCEMOEDNS-TEGYWKYG 117
DB 85 EYWDNRTRIYKDTAIFRVLNLTIRGYNQSETGSHNIQAMYGCDVCPDGLLGGFWQFG 144
QY 118 YDQDHLFCFDDTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQOLLELGR 177
DB 145 YDGRDYIALNEELRSWTAADTAQAQITKRKWEAAG-AAETWRNLEGCEVWLRRLYLENGK 203
QY 178 GVLDDQVPLVVKYTHH-VTSSTVTLRCALNYYQNTMVKDKQPMDAKEPEPKDVL 236
DB 204 DTLRLADPPKXVTHHSISDREVTLCRWALGFYPERISLTWQREGD-QTDMLVETRP 262
QY 237 NGDGTQGMITLAVPGGEQRYTCOVHPGLDPLIVIME 276
DB 263 SGGDTQKWAALVVPSEGEQRYTCRVQHEGLQEPILRW 302

RESULT 12
ID 1A11 HUMAN STANDARD; PRT; 365 AA.
AC P13746;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE HLA class I histocompatibility antigen, A-11 alpha chain precursor.
GN HLA-A OR HLAA.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A. (A*1101/A*1102).
RX MEDLINE=89030641; PubMed=2460344;
RA Mayer W.E., Jonker M., Klein D., Ivanyi P., van Seventer G.,
RA Klein J.;
RT "Nucleotide sequences of chimpanzee MHC class I alleles: evidence for
RT trans-species mode of evolution.";
RL EMBO J. 7:2765-2774(1988).
RN [2]
RP SEQUENCE FROM N.A. (A*1101/A*1102).
RX MEDLINE=94287401; PubMed=8016845;
RA Lin L., Tokunaga K., Ishikawa Y., Bannai M., Kashiwase K.,
RA Kuwata S., Akaza T., Tadokoro K., Shibata Y., Juji T.;
RT "Sequence analysis of serological HLA-A11 split antigens, A11.1 and
RT A11.2.";
RL Tissue Antigens 43:78-82(1994).
RN [3]
RP SEQUENCE OF 26-365 FROM N.A. (A*1101).
RX MEDLINE=87192928; PubMed=2437024;
RA Cowan E.P., Jelachich M.L., Bidison W.E., Coligan J.E.;
RT "DNA sequence of HLA-A11: remarkable homology with HLA-A3 allows
RT identification of residues involved in epitopes recognized by

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RT antibodies and T cells.";
RL Immunogenetics 25:241-250(1987).
CC -I- FUNCTION: Involved in the presentation of foreign antigens to
CC the immune system.
CC -I- SUBUNIT: Dimer of alpha chain and a beta chain (beta-2-
CC microglobulin).
CC -I- POLYMORPHISM: THE FOLLOWING ALLELES OF A-11 ARE KNOWN: A*1101 (A-
CC 11E) AND A*1102 (A-11K). THE SEQUENCE SHOWN IS THAT OF A*1101.
CC -----
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CC -----
DR EMBL; X13111; CAA31503.1; -
DR EMBL; X13112; CAA31504.1; -
DR EMBL; D16841; BAA04117.1; -
DR EMBL; D16842; BAA04118.1; -
DR EMBL; M16010; AAA65449.1; -
DR EMBL; M16007; AAA65449.1; JOINED.
DR EMBL; M16008; AAA65449.1; JOINED.
DR EMBL; M16009; AAA65449.1; JOINED.
DR PIR; I83063; I83063.
DR PIR; S03536; A47636.
DR HSP; O19673; 1HSB.
DR MIM; 142800; -
DR GO; GO:0005887; C:integral to plasma membrane; NAS.
DR GO; GO:0030106; F:MHC class I receptor activity; NAS.
DR GO; GO:0006955; P:immune response; NAS.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; IG_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGL1_1.
DR PROSITE; PS50835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal; Polymorphism.
FT SIGNAL 1 24
FT CHAIN 25 365 HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT A-11 ALPHA CHAIN.
FT DOMAIN 25 114 EXTRACELLULAR ALPHA-1.
FT DOMAIN 115 206 EXTRACELLULAR ALPHA-2.
FT DOMAIN 207 298 EXTRACELLULAR ALPHA-3.
FT DOMAIN 299 308 CONNECTING PEPTIDE.
FT TRANSMEM 309 332
FT DOMAIN 333 365 CYTOPLASMIC TAIL.
FT CARBOHYD 110 110 N-LINKED (GLCNAC. . .) (BY SIMILARITY).
FT DISULFID 125 188 BY SIMILARITY.
FT DISULFID 227 283 BY SIMILARITY.
FT VARIANT 43 43 E -> K (IN ALLELE A*1102).
FT SEQUENCE 365 AA; 40937 MW; FE449CE2D4BF6CC5 CRC64;

Query Match 33.8%; Score 514; DB 1; Length 365;
Best Local Similarity 39.4%; Pred. No. 7e-36;
Matches 109; Conservative 47; Mismatches 113; Indels 8; Gaps 7;

QY 5 SHSLHLYFMGASEQDGLSLFEALGVDDQLFVYDDB--SRVRPRTWVSSRISSQW 62
DB 26 SHSMRYFTTSVSRPGRGEPRPIAVGVYDDTQFVFRSDAASQRMPEAPWISQE-GPEYV 84
QY 63 LQLSOSLKGWDMFTVDFTWIMENHNHKSKE-SHTLQVILGCEMOEDNS-TEGYWKYG 120
DB 85 DQETNRVKAQSQDTRDVLGTLRGYTNQSDGGHTTQIMYGCDVCPDGLRGRDAYDG 144
QY 121 QDHLFCFDDTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQOLLELGR 180

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Db 145 KDVIALLNEDLSNTAADMAAQITKRWEAAH-AAEQRAYLEGRVCWEVRLRYLNGKLT 203
QY 181 DQOVPPPLVKVTH-VTSSVTTLCRALNYPQNTWKWLKDQPMADAKFEPKDVLPNGD 239
Db 204 QRTDPPKTHMTHTPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTLTVETRPAGD 262
QY 240 GTYQGITLAVPPEQRQYTCQVEHGLDQPLIVWE 276
Db 263 GTFQKAAVVVPSGEQRQYTCVQHEGLPKPLTLRWE 299

RESULT 13
ID 1A03 HUMAN STANDARD; PRT; 370 AA.
AC P04439;
DT 13-AUG-1987 (Rel. 05, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE HLA class I histocompatibility antigen, A-3 alpha chain precursor.
GN HLA-A OR HLAA.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN 1;
RP SEQUENCE FROM N.A. (A*0301).
RX MEDLINE=84207948; PubMed=6609814;
RA Strachan T., Sodoyer R., Damotte M., Jordan B.R.;
RT "Complete nucleotide sequence of a functional class I HLA gene,
RT HLA-A3: implications for the evolution of HLA genes.";
RL EMBO J. 3:887-894 (1984).
RN 2;
RP SEQUENCE FROM N.A. (A*0302).
RX MEDLINE=85290871; PubMed=2993417;
RA Cowan E.P., Jordan B.E., Colligan J.E.;
RT "Molecular cloning and DNA sequence analysis of genes encoding
RT cytotoxic T lymphocyte-defined HLA-A3 subtypes: the E1 subtype.";
RL J. Immunol. 135:2835-2841 (1985).
CC -1- FUNCTION: Involved in the presentation of foreign antigens to
CC the immune system.
CC -1- SUBUNIT: Dimer of alpha chain and a beta chain (beta-2-
CC microglobulin).
CC -1- POLYMORPHISM: THE FOLLOWING ALLELES OF A-3 ARE KNOWN: A*0301 (A-
CC 3.1) AND A*0302. THE SEQUENCE SHOWN IS THAT OF A*0301.
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CC -----
CC EMBL; X00492; CAA25162.1; ALT_TERM.
DR PIR; A02192; HLHUA3.
DR HSP; O19673; 1HSB.
DR STM; 142800;
DR GO; GO:0005887; C: integral to plasma membrane; NAS.
DR GO; GO:0030106; F: MHC class I receptor activity; NAS.
DR GO; GO:0006955; P: immune response; NAS.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_1.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGc1; 1.
DR PROSITE; PSS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW MHC I; Transmembrane; Glycoprotein; signal; Polymorphism.
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FT SIGNAL 1 29 HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT CHAIN 30 370 A-3 ALPHA CHAIN
FT DOMAIN 30 119 EXTRACELLULAR ALPHA-1.
FT DOMAIN 120 211 EXTRACELLULAR ALPHA-2.
FT DOMAIN 212 303 EXTRACELLULAR ALPHA-3.
FT DOMAIN 304 313 CONNECTING PEPTIDE.
FT TRANSMEM 314 337 CYTOPLASMIC TAIL.
FT DOMAIN 338 370 N-LINKED (GLUCNAC. .) (BY SIMILARITY).
FT CARBOHYD 115 115 BY SIMILARITY.
FT DISULFID 130 193 BY SIMILARITY.
FT DISULFID 232 288 E -> V (IN ALLELE A*0302).
FT VARIANT 181 181 /FTID=VAR 004351.
FT VARIANT 185 185 L -> Q (IN ALLELE A*0302).
FT /FTID=VAR 004352.
SQ SEQUENCE 370 AA; 41368 MW; ABBLFA77460318A2 CRC64;
Query Match 33.7%; Score 512; DB 1; Length 370;
Best Local Similarity 39.6%; Pred. No. 1.1e-35;
Matches 110; Conservative 47; Mismatches 111; Indels 10; Gaps 8;
QY 5 SHSLHYLFMGASEQDGLSLFALGVDDQLFVVDDE--SRVEPRTPWSSRSQW 62
Db 31 SHSMRYFTTSVSRPGRGEPRTIAGVYDDTQFVRFSDAASQRMPEAPWIEQ-GPEYW 89
QY 63 LQLSLSLKGWDHMTFVDFWFMENHNHNSKE-SHTLQVILGCEMQEDNS-TEGYWKYGDG 120
Db 90 DQETRNVAQAQSQTDRVDLGLTGLGYNQSEAGSHTIQIMYGCDVGSDFRLGTRQDAYDG 149
QY 121 QHLSFCPTDLWRAAEPRAPWTKLEWE--RHKIRARQNAYLERDCPAQLQELLELRGV 179
Db 150 KDVIALLNEDLSNTAADMAAQITKRWEAAH-AAEQRAYLDGTCVWELRYLNGKET 207
QY 180 LDOQVPPPLVKVTH-VTSSVTTLCRALNYPQNTWKWLKDQPMADAKFEPKDVLPNG 238
Db 208 LQRTDPPKTHMTHTPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTLTVETRPAG 266
QY 239 DQTYQGITLAVPPEQRQYTCQVEHGLDQPLIVWE 276
Db 267 DGTFOKAAVVVPSGEQRQYTCVQHEGLPKPLTLRWE 304

RESULT 14
ID 1A80 HUMAN STANDARD; PRT; 365 AA.
AC Q09160;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE HLA class I histocompatibility antigen, AW-80 (A-1) alpha chain
DE precursor.
GN HLA-A OR HLAA.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN 1;
RP SEQUENCE FROM N.A.
RX MEDLINE=94245293; PubMed=8188325;
RA Balas A., Garcia-Sanchez F., Gomez-Reino F., Vicario J.L.;
RT "Characterization of a new and highly distinguishable HLA-A allele in
RT a Spanish family.";
RL Immunogenetics 39:452-452 (1994).
RN 2;
RP SEQUENCE FROM N.A.
RA Domena J.D.;
RL Submitted (JUN-1993) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
CC THE IMMUNE SYSTEM.
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN).
CC -1- POLYMORPHISM: THE ONLY ALLELE OF AW-80 KNOWN IS A*8001 WHICH IS
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CC -----

DR EMBL; U03754; AAC04322.1; --
 DR EMBL; L18898; AAA17012.1; --
 DR PIR; I59638; I38439.
 DR HSP; Q95352; IHHK.
 DR MIM; I42800; --
 DR GO; GO:0005887; C: integral to plasma membrane; NAS.
 DR GO; GO:0030106; F: MHC class I receptor activity; NAS.
 DR GO; GO:0006955; P: immune response; NAS.
 DR InterPro; InterPro0110; Ig-like.
 DR InterPro; InterPro03597; Ig-cl.
 DR InterPro; InterPro03006; Ig_MHC.
 DR InterPro; InterPro01039; MHC_I.
 DR Pfam; PF00047; Ig; 1.
 DR Pfam; PF00129; MHC_I; 1.
 DR PRINTS; PR01638; MHCCLASSI.
 DR PRODOM; PD000050; MHC_I; 1.
 DR SMART; SM00407; IGL1; 1.
 DR PROSITE; PS00835; IG_LIKE; 1.
 DR PROSITE; PS00290; IG_MHC; 1.
 DR MHC_I; Transmembrane; Glycoprotein; Signal.
 DR SIGNAL 1 24
 DR CHAIN 25 365 HLA CLASS I HISTOCOMPATIBILITY ANTIGEN.
 DR FT
 DR FT DOMAIN 25 114 AW-80(A-1) ALPHA CHAIN.
 DR FT DOMAIN 115 206 EXTRACELLULAR ALPHA-1.
 DR FT DOMAIN 207 298 EXTRACELLULAR ALPHA-2.
 DR FT DOMAIN 299 308 EXTRACELLULAR ALPHA-3.
 DR FT TRANSMEM 309 332 CONNECTING PEPTIDE.
 DR FT DOMAIN 333 365 CYTOPLASMIC TAIL.
 DR FT CARBOHYD 110 110 N-LINKED (GLCNAC. . .) (BY SIMILARITY).
 DR FT DISULFID 125 188 BY SIMILARITY.
 DR FT DISULFID 227 283 BY SIMILARITY.
 DR SEQUENCE 365 AA; 40791 MW; CE1BC1CD60CA8FA8 CRC64;
 Query Match 33.6%; Score 510; DB 1; Length 365;
 Best Local Similarity 38.3%; Pred. No. 1.5e-35;
 Matches 106; Conservative 53; Mismatches 110; Indels 8; Gaps 7;
 QY 5 SHSLHYLFMGASQDGLSLFELGVYDDQLFFVYDDE--SRVERPTWSSRISSQW 62
 DB 26 SHSMRYFFTSVSRPGRGEPRFIAVGYYVDDSQFVQFSDAASQRMPEAPWIEQE-EPEYW 84
 QY 63 LQLSQSLKGDHMTFVDFWTIMENHNSKE-SHTLQVLGCENQEDNS-TEGYWKYGYDG 120
 DB 85 DEETRNVKAHSQNRNLANGLRGYYNQSDGSHTIQIMYGCVDGSGRFLRGYRDAYDG 144
 QY 121 QDHLFPCPTDLWRAAEPRAPWTKLEWRKIKRANQRNAYLRDCPAQLQQLLELGRGVL 180
 DB 145 KDVIALLNEDLRSTWADMAAQITKRWEAR-RAEQIRAYLEGECVDGLRRLYENKETYL 203
 QY 181 DQOVPLPVKVTTH-VTSSVTTLCRALNYPYQNTMKWLKQKQMDAKSEFPKDVLPNGD 239
 DB 204 QRTDPPKTHMTHPISDHEATLRCWALSFPASITLTWQDGDSD-QTQDTVELVETRPAGD 262
 QY 240 GTYQGITLAVPGEQRQYTCVQEHFGLDPLIVIE 276
 DB 263 GTFQKAAVVPVSGKSKRYTCHVOHGLPEPLRLWE 299
 RESULT 15
 ID 1A31_HUMAN STANDARD; PRT; 365 AA.
 AC F16189; Q98137; Q9TQ24;

DT 01-APR-1990 (Rel. 14, Created)
 DT 01-APR-1993 (Rel. 25, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE HLA class I histocompatibility antigen, A-31 alpha chain precursor
 DE (Aw-19).
 GN HLA-A OR HLA-A.
 OS Homo sapiens (Human).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 OC NCBI_TaxID=9606;
 OX [1]
 RN SEQUENCE FROM N.A. (A*3101).
 RX MEDLINE=90038496; PubMed=2478623;
 RA Kato K., Trapani J.A., Allopenna J., Dupont B., Yang S.Y.;
 RT "Molecular analysis of the serologically defined HLA-Aw19 antigens. A
 RT genetically distinct family of HLA-A antigens comprising A29, A31,
 RT A32, and A*33, but probably not A30.";
 RL J. Immunol. 143:3371-3378(1989).
 RN [2]
 RN SEQUENCE FROM N.A. (A*3101).
 RX MEDLINE=92269955; PubMed=1317015;
 RA Belich M.P., Madrigal J.A., Hildebrand W.H., Zemmour J.,
 RA Williams R.C., Luz R., Petzl-Erler M.L., Parham P.;
 RT "Unusual HLA-B alleles in two tribes of Brazilian Indians.";
 RL Nature 357:326-329(1992).
 RN [3]
 RN SEQUENCE FROM N.A. (A*3101).
 RX MEDLINE=96387675; PubMed=8795145;
 RA Arnett K.L., Adams E.J., Parham P.;
 RT "On the sequence of A*3101.";
 RL Tissue Antigens 47:428-430(1996).
 RN [4]
 RN SEQUENCE OF 9-365 FROM N.A. (A*3101).
 RX MEDLINE=92269956; PubMed=1599035;
 RA Watkins D.I., McAdam S.N., Liu X., Stang C.R., Milford E.L.,
 RA Levine C.G., Garber T.L., Dogan A.L., Lord C.I., Ghim S.H.,
 RA Troup G.M., Hughes A.L., Letvin N.L.;
 RT "New recombinant HLA-B alleles in a tribe of South American
 RT Amerindians indicate rapid evolution of MHC class I loci.";
 RL Nature 357:329-333(1992).
 RN [5]
 RN SEQUENCE FROM N.A. (A*3104).
 RA Bettinotti M.P., Dhillion G., Hackett J., Simonis T.B., Marincola F.M.;
 RT "A New HLA-A*31 allele.";
 RL Submitted (MAY-1999) to the EMBL/GenBank/DBJ databases.
 RN [6]
 RN SEQUENCE OF 26-206 FROM N.A. (A*3104).
 RA Mitsushige Y.;
 RT "New HLA-A31 allele identified in African American population.";
 RL Submitted (NOV-1998) to the EMBL/GenBank/DBJ databases.
 CC -!- FUNCTION: Involved in the presentation of foreign antigens to
 CC the immune system.
 CC -!- SUBUNIT: Dimer of alpha chain and a beta chain (beta-2-
 CC microglobulin).
 CC -!- POLYMORPHISM: THE FOLLOWING ALLELES OF A-31 ARE KNOWN: A*3101 AND
 CC A*3104. THE SEQUENCE SHOWN IS THAT OF A*3101.
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CC -----

DR EMBL; M30578; AAA59613.1; --
 DR EMBL; M84375; AAA59599.1; --
 DR EMBL; L78918; AAB05976.1; --
 DR EMBL; AF148863; AAD39981.1; --
 DR EMBL; AF105028; AAC79721.1; --
 DR EMBL; AF105027; AAC79721.1; JOINED.
 DR FIR; I72170; I72170.
 DR HSP; O19673; IHSB.

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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:05:29 ; Search time 32 Seconds
(without alignments)

2225.704 Million cell updates/sec

Title: US-10-092-404-2

Perfect score: 1520

Sequence: 1 RLRSLSLHFLMGASEQDL.....RYTCQVEHPGLDQPLIVME 276

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database :

SPTREMBL_23:*

1: sp_archaea:*

2: sp_bacteria:*

3: sp_fungi:*

4: sp_human:*

5: sp_invertebrate:*

6: sp_mammal:*

7: sp_mhc:*

8: sp_organelle:*

9: sp_phase:*

10: sp_plant:*

11: sp_rodent:*

12: sp_virus:*

13: sp_vertebrate:*

14: sp_unclassified:*

15: sp_rvirus:*

16: sp_bacteriap:*

17: sp_archaeap:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1140	75.0	358	11 Q8C2A6	Q8C2A6 mus musculus
2	1140	75.0	359	11 Q9D754	Q9D754 mus musculus
3	802	52.8	272	11 Q9R105	Q9R105 rattus norv
4	592	38.9	116	4 Q9HC69	Q9HC69 homo sapien
5	547.5	36.0	359	7 Q8HX81	Q8HX81 ornithorhyn
6	543.5	35.8	340	7 Q9BD50	Q9BD50 pongo pygma
7	542.5	35.7	334	7 Q9TOK3	Q9TOK3 homo sapien
8	542.5	35.7	341	4 Q9NPL2	Q9NPL2 homo sapien
9	542.5	35.7	341	7 Q9S460	Q9S460 homo sapien
10	542.5	35.7	341	7 Q9BCU3	Q9BCU3 pan troglod
11	540.5	35.6	354	7 Q9SHB3	Q9SHB3 anas platyr
12	539.5	35.5	341	7 Q9BCU4	Q9BCU4 pan troglod
13	530	34.9	105	4 Q9HC71	Q9HC71 homo sapien
14	521	34.3	356	7 Q8HX66	Q8HX66 sus scrofa
15	520	34.2	332	7 Q30990	Q30990 pan troglod
16	520	34.2	365	7 Q9TLP7	Q9TLP7 pan troglod

17	518	34.1	365	7	Q9TQP6	Q9TQP6 homo sapien
18	518	34.1	371	7	Q9TQP7	Q9TQP7 homo sapien
19	517	34.0	352	7	Q8MHT1	Q8MHT1 sus scrofa
20	517	34.0	364	7	O19243	O19243 sus scrofa
21	517	34.0	365	7	Q9MYG4	Q9MYG4 homo sapien
22	515	33.9	365	7	Q29747	Q29747 homo sapien
23	514	33.8	273	7	Q9S1G6	Q9S1G6 homo sapien
24	514	33.8	352	7	Q8SPA9	Q8SPA9 sus scrofa
25	514	33.8	361	7	Q8HX63	Q8HX63 sus scrofa
26	514	33.8	364	7	Q8HX61	Q8HX61 sus scrofa
27	514	33.8	365	7	Q9MYI5	Q9MYI5 homo sapien
28	513	33.8	330	7	O19356	O19356 macaca mula
29	513	33.8	331	7	O02944	O02944 macaca mula
30	513	33.8	333	7	Q98030	Q98030 papio anubi
31	513	33.8	333	7	Q98031	Q98031 papio anubi
32	512	33.7	129	4	Q9UX37	Q9UX37 homo sapien
33	512	33.7	330	7	O02947	O02947 macaca mula
34	512	33.7	330	7	O02946	O02946 macaca mula
35	512	33.7	365	7	O19756	O19756 homo sapien
36	511	33.6	331	7	O02945	O02945 macaca mula
37	511	33.6	357	7	Q30886	Q30886 pan paniscu
38	511	33.6	363	7	Q9MXI5	Q9MXI5 pan troglod
39	511	33.6	363	7	Q9MWK4	Q9MWK4 gorilla gor
40	511	33.6	365	7	Q9MXI6	Q9MXI6 pan troglod
41	511	33.6	365	7	Q9MXM7	Q9MXM7 pan troglod
42	510	33.6	360	7	Q9GJ24	Q9GJ24 homo sapien
43	510	33.6	364	7	Q8SPA4	Q8SPA4 sus scrofa
44	510	33.6	365	7	Q30900	Q30900 pan paniscu
45	510	33.6	365	7	Q9BCN0	Q9BCN0 homo sapien

ALIGNMENTS

RESULT 1

Q8C2A6 PRELIMINARY; PRT; 358 AA.
ID Q8C2A6
AC Q8C2A6;
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DB Hemochromatosis.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=NOD; TISSUE=Thymus;
RX MEDLINE=22354683; PubMed=12466851;
RA The FANTOM Consortium.
RA the RIKEN Genome Exploration Research Group Phase I & II Team;
RT "Analysis of the mouse transcriptome based on functional annotation of
RT 60,770 full-length cDNAs.";
RL Nature 420:563-573(2002).
DR EMBL; AK089886; BAC40688.1; -.
SQ SEQUENCE 358 AA; 40421 MW; EE88FB6E5AAC844D CRC64;

Query Match 75.0%; Score 1140; DB 11; Length 358;
Best Local Similarity 72.2%; Pred. No. 2.5e-99;
Matches 203; Conservative 30; Mismatches 40; Indels 8; Gaps 1;
QY 4 RSHSLHFLMGASEQDLGLSFEALGYDDQLFVFYDDSSRRVPTPWSSRISSQWL 63
Db 29 RSHSLHFLMGASEQDLGLSFEALGYDDQLFVFYDDSSRRVPTPWSSRISSQWL 89
QY 64 QLSQSLKGDHMTFTVDFTWIMENHNSK-----ESHTLQVILGCENQENSTGYMK 115
Db 89 HLSQSLKGDHMTFTVDFTWIMENHNSK-----ESHTLQVILGCENQENSTGYMK 148
QY 116 YGVDGDHLEFPCDTLDWRAAEFRAPWTKLEWERHKIRAKONRAYLERCPQLQLEL 175
Db 149 YGVDGDHLEFPCDTLDWRAAEFRAPWTKLEWERHKIRAKONRAYLERCPQLQLEL 208

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QY 176 GRGVLQVPLVKVTHVTSVTLRCALNYYPONITMKLQKQPMDAKEFEFKDVL 235
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 209 GRGVLQVPLVKVTHVTSVTLRCALNYYPONITMKLQKQPMDAKEFEFKDVL 268
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
QY 236 PNGDGTQYQWITLAVPPGEQRYTCQVEHPGLDQPLVIWE 276
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 269 PNGDGTQYQWITLAVPPGEQRYTCQVEHPGLDQPLVIWE 309
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:

RESULT 2
Q9D754 PRELIMINARY; PRT; 359 AA.
AC Q9D754;
DT 01-JUN-2001 (T-EMBLrel. 17, Created)
DT 01-JUN-2001 (T-EMBLrel. 17, Last sequence update)
DT 01-MAR-2003 (T-EMBLrel. 23, Last annotation update)
DE Adult male tongue cDNA, RIKEN full-length enriched library,
DE clone:2310032M04, full insert sequence.
GN HFE.
OS Mus musculus (Mouse).
OC Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RP SEQUENCE FROM N.A.
RC SPRAIN=C57BL/6J; TISSUE=Tongue;
RX MEDLINE=2108560; PubMed=11217851;
RA Kawai J., Shingawa A., Shibata K., Yoshino M., Itoh M., Ishii Y.,
RA Arakawa T., Hara A., Fukunishi Y., Konno H., Adachi J., Fukuda S.,
RA Aizawa K., Izawa M., Nishi K., Kiyosawa H., Kondo S., Yamanaka I.,
RA Saito T., Okazaki Y., Gojobori T., Bono H., Kasukawa T., Saito R.,
RA Kadota K., Matsuda H.A., Ashburner M., Batalov S., Casavant T.,
RA Fleischmann W., Gaasterland T., Giesi C., King B., Kochiwa H.,
RA Kuehl P., Lewis S., Matsuo Y., Nikaide I., Pesole G., Quackenbush J.,
RA Schram L.M., Staubli F., Suzuki R., Tomita M., Wagner L., Washio T.,
RA Sakai K., Okido T., Furuno M., Aono H., Baldarelli R., Barsh G.,
RA Blake J., Boffelli D., Bojunga N., Carninci P., de Bonaldo M.F.,
RA Brownstein M.J., Bult C., Fletcher C., Fujita M., Gariboldi M.,
RA Guncincich S., Hill D., Hofmann M., Hume D.A., Kamiya M., Lee N.H.,
RA Lyons P., Marchionni L., Mashima J., Mazzarelli J., Mombaerts P.,
RA Nordone P., Ring B., Ringwald M., Rodriguez I., Sakamoto N.,
RA Suzuki H., Sato K., Schoenbach C., Seya T., Shibata Y., Storch K.-F.,
RA Suzuki H., Toyo-oka K., Wang K.H., Weitz C., Whittaker C., Wilming L.,
RA Wyshaw-Boris A., Yoshida K., Hasegawa Y., Kawaji H., Kohtsuki S.,
RA Havaehizaki Y.
RT "Functional annotation of a full-length mouse cDNA collection.";
RC Nature 409:685-690(2001).
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
CC ENBL; AK009581; BAB26373.1; -.
DR HSSP; Q30201; 1A6Z.
DR MGD; MGI:109191; Hfe.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR PRINTS; PR01638; MHC_I.
DR PROSITE; PS00050; MHC_I.
DR PROSITE; PS00290; Ig_MHC; 1.
DR GlycoProtein; Transmembrane.
KW Glycoprotein; Transmembrane.
SQ SEQUENCE 359 AA; 40534 MW; 586657B7F9FF20B4 CRC64;

Query Match 75.0%; Score 1140; DB 11; Length 359;
Best Local Similarity 72.2%; Pred. No. 2.5e-99;
Matches 203; Conservative 30; Mismatches 40; Indels 8; Gaps 1;

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QY 4 RSHSLHYLFMGASEQDLGLSFEALGYVDDQLFVYVDDSRVRPRTPWVSRISSQML 63
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 30 RSHSLHYLFMGASEQDLGLSFEALGYVDDQLFVYVDDSRVRPRTPWVSRISSQML 89
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
QY 64 QLSQSILKGDHMTFTDFTIMENHNHSHK-----ESHTLVILGCEMOEDNSTGYWK 115
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 90 HLSQSLKGDHMTFTDFTIMGNHNSKVTGLGVVSESHILQVVLGCEVHEDNSTSGFWR 149
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
QY 116 YGYDGDHLEFCFPTDLDWRAAPRAWPTKLEWERHKIRARQNRVYLERDCPAQLQELLE 175
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 150 YGYDGDHLEFCFPTLNWSAEPGAWATKVEWDEHKIRAKQNRDYLEKDCPEQLKELLE 209
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
QY 176 GRGVLQVPLVKVTHVTSVTLRCALNYYPONITMKLQKQPMDAKEFEFKDVL 235
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 210 GRGVLQVPLVKVTHVTSVTLRCALNYYPONITMKLQKQPMDAKEFEFKDVL 269
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
QY 236 PNGDGTQYQWITLAVPPGEQRYTCQVEHPGLDQPLVIWE 276
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 270 PNGDGTQYQWITLAVPPGEQRYTCQVEHPGLDQPLVIWE 310
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:

RESULT 3
Q9RI05 PRELIMINARY; PRT; 272 AA.
AC Q9RI05;
DT 01-MAY-2000 (T-EMBLrel. 13, Created)
DT 01-MAY-2000 (T-EMBLrel. 13, Last sequence update)
DT 01-MAR-2003 (T-EMBLrel. 23, Last annotation update)
DE Hemochromatosis gene product HFE splice variant delE2.
OS Rattus norvegicus (Rat).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
OX NCBI_TaxID=10116;
RN [1]
RP SEQUENCE FROM N.A.
RC STRAIN=Wistar; TISSUE=Testis;
RA Liew Y.-F., Shaw N.-S.;
RT "Alternative splice variant of the hemochromatosis gene HFE in iron
RT overloaded rats.";
RL Submitted (AUG-1999) to the EMBL/GenBank/DBJ databases.
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
CC ENBL; AF176534; AAD49965.1; -.
DR HSSP; Q30201; 1A6Z.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR PRINTS; PR01638; MHC_I.
DR PROSITE; PS00050; MHC_I.
DR PROSITE; PS00407; IGL1; 1.
DR PROSITE; PS00835; IG LIKE; 1.
DR PROSITE; PS00290; Ig_MHC; 1.
DR GlycoProtein; Transmembrane.
KW Glycoprotein; Transmembrane.
SQ SEQUENCE 272 AA; 30757 MW; 1D91063CCBEFF5502 CRC64;

Query Match 52.8%; Score 802; DB 11; Length 272;
Best Local Similarity 75.1%; Pred. No. 1.4e-67;
Matches 139; Conservative 22; Mismatches 24; Indels 0; Gaps 0;

QY 92 ESHTLVILGCEMOEDNSTGYWKYDGDHLEFCFPTDLDWRAAPRAWPTKLEWERHK 151
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 39 ESHILQVILGCEVHEDNSTSGFWKYGVDGDHLEFCFPTLNWSAEPRAWATKWEHR 98
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
QY 152 IRARQNRVYLERDCPAQLQELLEGRVLDQVPLVKVTHVTSVTLRCALNYYPQ 211
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
Db 99 IRARQNRVYLERDCPAQLQELLEGRVLDQVPLVKVTHVTSVTLRCALNYYPQ 158
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:
QY 212 NITMKWLKQPMDAKEFEFKDVL PNGDGTQYQWITLAVPPGEQRYTCQVEHPGLDQPL 271
    |||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:|||||:

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Db 83 RYTQLRGQQMFVKELKRLQRHYNHS -GSHTYORMIGCELLBGSTTGFLQAYDQDF 141
Qy 124 LEPCPTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVLDQ 183
Db 142 LIFNKDTLSLWADVNVHTIKRANEQHELQYQKWLKEECIAWLKRFLEYGKDTLQRT 201
Qy 184 VPPLVKVTHVHT-SSVTTLCRALNYYPQNIITMKWLKDKQPMDAKEFEFPKQVLPNGDGT 242
Db 202 EPPLVRVNRKETFPFGVTTALFCKAHGFYPPEIYMTWMKGEEI-VQMDYDGLILPSGDGT 260
Qy 243 QGWITLAVPPGEEQRYTCQVHPGLDQPLIV 273
Db 261 QTWASFELDPQSSNLYSCHVEHCGVHMVLQV 291

RESULT 7

Q9TK3 ID Q9TK3 PRELIMINARY; PRT; 334 AA.
AC Q9TK3
DT 01-MAY-2000 (TREMBLrel. 13, Created)
DT 01-MAY-2000 (TREMBLrel. 13, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE MHC class I-related protein MRL (Fragment).
GN MRL.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Placenta;
RX MEDLINE=99003494; PubMed=9784382;
RA Yamaguchi H., Kuroawa Y., Hashimoto K.;
RT "Expanded genomic organization of conserved mammalian MHC class I-
RT related genes, human MRL and its murine ortholog.";
RL Biochem. Biophys. Res. Commun. 250:558-564 (1998).
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
DR EMBL; AF073485; AAC72900.1; JOINED.
DR EMBL; AF073484; AAC72900.1; JOINED.
DR HSSP; Q30201; IA6Z.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC I; 1.
DR ProDom; PD000050; MHC I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PS00835; IG LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; Transmembrane.
FT NON TER 1
SQ SEQUENCE 334 AA; 38586 MW; 4C3E8A248A39BA4 CRC64;

Query Match 35.7%; Score 542.5; DB 7; Length 334;
Best Local Similarity 39.5%; Pred. No. 6e-43;
Matches 107; Conservative 50; Mismatches 111; Indels 3; Gaps 3;
Qy 4 RSHSLYLFMGASEQDLGLSLFALGYVDQDLFFVYDDSRREPRTPWSSRISSQMWL 63
Db 16 RTHSLRYFLGVSDPIHGVPFISGVYDSDHPITTYDSVTRQKEPRAPMAENLADPHE 75
Qy 64 QLSQSLKGWDMFTVDFWTIMENHNHKSHTLQVLGCEMQEDNSTEGYWKYGYDQDH 123
Db 76 RYTQLRGQQMFVKELKRLQRHYNHS -GSHTYORMIGCELLBGSTTGFLQAYDQDF 134
Qy 124 LEPCPTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVLDQ 183
Db 135 LIFNKDTLSLWADVNVHTIKRANEQHELQYQKWLKEECIAWLKRFLEYGKDTLQRT 194

Qy 184 VPPLVKVTHVHT-SSVTTLCRALNYYPQNIITMKWLKDKQPMDAKEFEFPKQVLPNGDGT 242
Db 195 EPPLVRVNRKETFPFGVTTALFCKAHGFYPPEIYMTWMKGEEI-VQMDYDGLILPSGDGT 253
Qy 243 QGWITLAVPPGEEQRYTCQVHPGLDQPLIV 273
Db 254 QWASIELDPQSSNLYSCHVEHCGVHMVLQV 284

RESULT 8

Q9NPL2 ID Q9NPL2 PRELIMINARY; PRT; 341 AA.
AC Q9NPL2
DT 01-OCT-2000 (TREMBLrel. 15, Created)
DT 01-OCT-2000 (TREMBLrel. 15, Last sequence update)
DT 01-MAR-2003 (TREMBLrel. 23, Last annotation update)
DE MRL protein.
GN MRL.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=Peripheral blood;
RX MEDLINE=20470599; PubMed=11019920;
RA Parra-Cuadrado J.F., Navarro P., Mirones I., Setien F., Oteo M.,
RA Martinez-Naves E.;
RT "A study on the polymorphism of human MHC class I-related MRL gene and
RT identification of an MRL-like pseudogene.";
RL Tissue Antigens 56:170-172 (2000).
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
DR EMBL; AJ249778; CAB77667.1; -.
DR HSSP; Q30201; IA6Z.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC I; 1.
DR ProDom; PD000050; MHC I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PS00835; IG LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; Transmembrane.
SQ SEQUENCE 341 AA; 39366 MW; 2990C1F3F0A1CAD9 CRC64;

Query Match 35.7%; Score 542.5; DB 4; Length 341;
Best Local Similarity 39.5%; Pred. No. 6.2e-43;
Matches 107; Conservative 50; Mismatches 111; Indels 3; Gaps 3;
Qy 4 RSHSLYLFMGASEQDLGLSLFALGYVDQDLFFVYDDSRREPRTPWSSRISSQMWL 63
Db 23 RTHSLRYFLGVSDPIHGVPFISGVYDSDHPITTYDSVTRQKEPRAPMAENLADPHE 82
Qy 64 QLSQSLKGWDMFTVDFWTIMENHNHKSHTLQVLGCEMQEDNSTEGYWKYGYDQDH 123
Db 83 RYTQLRGQQMFVKELKRLQRHYNHS -GSHTYORMIGCELLBGSTTGFLQAYDQDF 141
Qy 124 LEPCPTLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVLDQ 183
Db 142 LIFNKDTLSLWADVNVHTIKRANEQHELQYQKWLKEECIAWLKRFLEYGKDTLQRT 201
Qy 184 VPPLVKVTHVHT-SSVTTLCRALNYYPQNIITMKWLKDKQPMDAKEFEFPKQVLPNGDGT 242
Db 202 EPPLVRVNRKETFPFGVTTALFCKAHGFYPPEIYMTWMKGEEI-VQMDYDGLILPSGDGT 260
Qy 243 QGWITLAVPPGEEQRYTCQVHPGLDQPLIV 273

455


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Matches 94; Conservative 3; Mismatches 3; Indels 0; Gaps 0;
QY 89 HSKESHTLQVILGCEMEDNSTGYKYGVDGQHLFCFPTLDWRAAEPRAPWTKLEWE 148
DB 1 HTKESHTLQVILGCEMEDNSTGYKYGVDGQHLFCFPTLDWRAAEPRAPWTKLEWE 60
QY 149 RHKIRARONRAYLERDCPAQLQQLLELGRGVLDQVPPV 188
DB 61 GHKVRARONGAYLERDCPAQLQQLLELGRGVLDQVPPV 100

RESULT 14
Q8HX66 PRELIMINARY; PRT; 356 AA.
AC Q8HX66;
DT 01-MAR-2003 (TrEMBLrel. 23, Created)
DT 01-MAR-2003 (TrEMBLrel. 23, Last sequence update)
DE MHC class I antigen (Fragment).
GN SLA-1.
OS Sus scrofa (Pig).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
NCBI_TaxID=9823;
RN [1]
RP SEQUENCE FROM N.A.
RA Martens G.W., Baker J.E., Smith D.M.;
RL Submitted (JUL-2002) to the EMBL/GenBank/DBJ databases.
DR EMBL; AY135589; AN35107.1; --
FT NON TER 1
SQ SEQUENCE 356 AA; 3585 MW; 94FC7A461DBF555B CRC64;

Query Match 34.3%; Score 521; DB 7; Length 356;
Best Local Similarity 39.9%; Pred. No. 7e-41;
Matches 110; Conservative 48; Mismatches 110; Indels 8; Gaps 7;
QY 6 HSLHYLFMGASEODLGLSLFEALGYVDDQLFVYDDE--SRVPEPTPWSSRISQMWL 63
DB 19 HSLRYFYTAVRPDLGDSRFIAGVYDDTQVFRFSDAPNPRMEPRAPWTKQE--GOEYWD 77
QY 64 QLSQSLKGWDHMTVDFTWIMENHNHKS--SHTLQVILGCEMEDN--STEGYKYGVDG 121
DB 78 EETRNANGSQNDVRDLKTLRGYNGSEAGSHTIQRMVCDVPGDGLLGRYDQDAYDGA 137
QY 122 DHLEFCFPTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVLD 181
DB 138 DYIALNEDLRSWTAADTAAGITTKRWEAANV-AEQERSYLEGTCVEWLQYLEMGKDTLQ 196
QY 182 QVPPPLVKVTHHTSSV--TTLRCALNYYPONTIMKWLKDKQPMDAKEFEPPKDVLPNGDG 240
DB 197 RAEPKTHVTRHPSSDLGVTRCWLALGFYKPEISLTWQREGD--QSQDMELVETRPSPGDG 255
QY 241 TYQGWTITLAVPPGEEQRYTCQVHPGLDQPLVIWE 276
DB 256 TFQKWAALVVPGEQSYTCHVQHEGLPKPLTLRWE 291

RESULT 15
Q30990 PRELIMINARY; PRT; 332 AA.
AC Q30990;
DT 01-NOV-1996 (TrEMBLrel. 01, Created)
DT 01-NOV-1996 (TrEMBLrel. 01, Last sequence update)
DE 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Chimpanzee MHC class I Chla chain (Fragment).
OS Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
NCBI_TaxID=9598;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=89235215; PubMed=2715640;
RA Farham P., Lawlor D.A., Lomen C.E., Ennis P.D.;

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RT "Diversity and diversification of HLA-A,B,C alleles.";
RL J. Immunol. 142:3937-3950(1989).
CC -I- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -I- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICRoglobulin) (BY SIMILARITY).
DR EMBL; M24047; AAA35426.1; --
DR HSSP; Q95352; IHHK.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig_c1.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IgC1; 1.
DR PROSITE; PS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; Transmembrane.
FT NON TER 332
SQ SEQUENCE 332 AA; 37433 MW; 9AA9A55DF9E79360 CRC64;

Query Match 34.2%; Score 520; DB 7; Length 332;
Best Local Similarity 40.1%; Pred. No. 7.9e-41;
Matches 111; Conservative 44; Mismatches 114; Indels 8; Gaps 7;
QY 5 HSLHYLFMGASEODLGLSLFEALGYVDDQLFVYDDE--SRVPEPTPWSSRISQMW 62
DB 26 HSMRYFFTSVSRPGEGEPRFIAGVYDDTQVFRFSDAASQRMPEPRAPWTKQE--GPEYW 84
QY 63 LQLSOSLKGWDHMTVDFTWIMENHNHKS--SHTLQVILGCEMEDNS--TEGYKYGVDG 120
DB 85 DQETRSAAHSQTDKVDLGLTGLRGYNGSEDSHTTIQIMYGVCDVSGDGRFLRGYRDAYDG 144
QY 121 QHLEFCFPTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVLD 180
DB 145 KYIYALNEDLRSWTAADTAAGITTKRWEAAH-AAEQRAYLEGTCVEWLRLYLENGKETL 203
QY 181 DOQVPLVKVTHH--VTSSVTLRCALNYYPONTIMKWLKDKQPMDAKEFEPPKDVLPNGD 239
DB 204 QRTDPPKTHHTHPISDHEATLRCWALGFYPAEITLTWQREGD--QTQDTLVEPTRPADG 262
QY 240 GYQGWITLAVPPGEEQRYTCQVHPGLDQPLVIWE 276
DB 263 GTFQKWAALVVPGEQRYTCHVQHEGLPKPLTLRWE 299

Search completed: August 5, 2003, 13:10:00
Job time : 34 secs

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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:10:04 ; Search time 33 Seconds
(without alignments)
993.264 Million cell updates/sec

Title: US-10-092-404-2
Perfect score: 1520
Sequence: 1 RLRSLSLHLYFMGASEQDL.....RYTCQVEHGLDPLIVIME 276

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 451899 seqs, 118759770 residues

Total number of hits satisfying chosen parameters: 451899

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications AA:

- 1: /cgn2_6/ptodata/1/pubpaa/US07_PUBCOMB.pep.*
- 2: /cgn2_6/ptodata/1/pubpaa/PCT_NEW_PUB.pep.*
- 3: /cgn2_6/ptodata/1/pubpaa/US06_NEW_PUB.pep.*
- 4: /cgn2_6/ptodata/1/pubpaa/US06_PUBCOMB.pep.*
- 5: /cgn2_6/ptodata/1/pubpaa/US07_NEW_PUB.pep.*
- 6: /cgn2_6/ptodata/1/pubpaa/PCTUS_PUBCOMB.pep.*
- 7: /cgn2_6/ptodata/1/pubpaa/US08_NEW_PUB.pep.*
- 8: /cgn2_6/ptodata/1/pubpaa/US08_PUBCOMB.pep.*
- 9: /cgn2_6/ptodata/1/pubpaa/US09A_PUBCOMB.pep.*
- 10: /cgn2_6/ptodata/1/pubpaa/US09B_PUBCOMB.pep.*
- 11: /cgn2_6/ptodata/1/pubpaa/US09C_PUBCOMB.pep.*
- 12: /cgn2_6/ptodata/1/pubpaa/US09_NEW_PUB.pep.*
- 13: /cgn2_6/ptodata/1/pubpaa/US10A_PUBCOMB.pep.*
- 14: /cgn2_6/ptodata/1/pubpaa/US10B_PUBCOMB.pep.*
- 15: /cgn2_6/ptodata/1/pubpaa/US10C_PUBCOMB.pep.*
- 16: /cgn2_6/ptodata/1/pubpaa/US10_NEW_PUB.pep.*
- 17: /cgn2_6/ptodata/1/pubpaa/US60_NEW_PUB.pep.*
- 18: /cgn2_6/ptodata/1/pubpaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1520	100.0	276	15	US-10-092-404-2
2	1513	99.5	276	15	US-10-092-404-1
3	1513	99.5	348	12	US-09-981-606-2
4	1493	98.2	276	15	US-10-092-404-3
5	514	33.8	92	14	US-10-016-634A-120
6	506	33.3	280	15	US-10-073-300-6
7	506	33.3	415	15	US-10-073-300-5
8	492	32.4	298	15	US-10-205-823-40
9	492	32.4	298	15	US-10-205-823-42
10	492	32.4	298	15	US-10-177-293-23
11	477	31.4	542	15	US-10-015-535-32
12	477	31.4	542	15	US-10-015-535-34
13	475	31.2	542	15	US-10-015-535-36
14	474	31.2	540	15	US-10-015-535-22
15	474	31.2	541	15	US-10-015-535-28

16	474	31.2	542	15	US-10-015-535-24	Sequence 24, Appl
17	474	31.2	542	15	US-10-015-535-26	Sequence 26, Appl
18	448	29.5	332	9	US-09-870-521-3	Sequence 3, Appli
19	445	29.3	540	15	US-10-015-535-30	Sequence 30, Appli
20	444	29.2	334	9	US-09-870-521-4	Sequence 4, Appli
21	358.5	23.6	170	9	US-09-925-301-1307	Sequence 1307, Ap
22	336	22.1	271	9	US-09-925-301-1431	Sequence 1431, Ap
23	279	18.4	181	11	US-09-013-077A-13	Sequence 13, Appl
24	275	18.1	145	9	US-09-810-560-8	Sequence 8, Appli
25	243	16.0	184	10	US-09-858-580-21	Sequence 21, Appl
26	243	16.0	184	11	US-09-847-172-21	Sequence 21, Appl
27	226	14.9	91	9	US-09-864-761-38005	Sequence 38005, A
28	223.5	14.7	171	15	US-10-144-929-116	Sequence 116, App
29	223	14.7	91	9	US-09-864-761-35461	Sequence 35461, A
30	210.5	13.8	104	9	US-09-925-302-835	Sequence 835, App
31	208.5	13.7	183	15	US-10-036-542-62	Sequence 62, Appli
32	207	13.6	117	9	US-09-810-560-9	Sequence 9, Appli
33	196.5	12.9	93	9	US-09-864-761-39479	Sequence 39479, A
34	196.5	12.9	110	9	US-09-864-761-35339	Sequence 35339, A
35	196.5	12.9	114	9	US-09-864-761-37988	Sequence 37988, A
36	174.5	11.5	261	10	US-09-925-664-30	Sequence 30, Appl
37	174	11.4	411	14	US-10-015-536-17	Sequence 17, Appl
38	173	11.4	110	10	US-09-796-692-799	Sequence 799, App
39	173	11.4	110	10	US-09-796-692-2139	Sequence 2139, Ap
40	173	11.4	110	15	US-10-040-862-799	Sequence 799, App
41	173	11.4	110	15	US-10-040-862-2139	Sequence 2139, Ap
42	171.5	11.3	285	10	US-09-756-983-24	Sequence 24, Appl
43	167	11.0	772	9	US-09-815-837-74	Sequence 74, Appl
44	166.5	11.0	448	14	US-10-081-281-111	Sequence 111, App
45	166	10.9	246	9	US-09-989-722-225	Sequence 225, App

ALIGNMENTS

RESULT 1

US-10-092-404-2
; Sequence 2, Application US/10092404
; Publication No. US20030073627A1

; GENERAL INFORMATION:

; APPLICANT: Feder, John N.
; Bjorkman, Pamela J.
; Schatzman, Randall C.

; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
; DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
; AND IRON DEFICIENCY DISEASES

; NUMBER OF SEQUENCES: 5

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Pennie & Edmonds, LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: NY

; COUNTRY: USA

; ZIP: 10036-2811

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Diskette

; OPERATING SYSTEM: Windows

; SOFTWARE: FastSeq for Windows Version 2.0b

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/10/092,404

; FILING DATE: 04-Mar-2002

; CLASSIFICATION: <Unknown>

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: US/09/094,964

; FILING DATE: June 12, 1998

; APPLICATION NUMBER: 08/876,010

; FILING DATE: June 13, 1997

; ATTORNEY/AGENT INFORMATION:

; NAME: Poissant, Brian M

; REGISTRATION NUMBER: 28,462

; REFERENCE/DOCKET NUMBER: 8907-0074-999

; TELECOMMUNICATION INFORMATION:

TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEFAX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 276 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
SEQUENCE DESCRIPTION: SEQ ID NO: 2:
US-10-092-404-2

Query Match 100.0%; Score 1520; DB 15; Length 276;
Best Local Similarity 100.0%; Pred. No. 1.7e-146;
Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISSQ 60
DB 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISSQ 60
QY 61 MWLQLSQSLKGDHMTVDFTWMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 61 MWLQLSQSLKGDHMTVDFTWMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
QY 121 QDHLFCPDTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 121 QDHLFCPDTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
QY 181 DQVPLVAVKTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
DB 181 DQVPLVAVKTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
DB 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276

RESULT 2

US-10-092-404-1
Sequence 1, Application US/10092404
Publication No. US20030073627A1
GENERAL INFORMATION:
APPLICANT: Feder, John N.
Bjorkman, Pamela J.
Schatzman, Randall C.
TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
AND IRON DEFICIENCY DISEASES
NUMBER OF SEQUENCES: 5
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds, LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10036-2811
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: Windows
SOFTWARE: FastSeq for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/10/092,404
FILING DATE: 04-Mar-2002
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/09/094,964
FILING DATE: June 12, 1998
APPLICATION NUMBER: 08/876,010
FILING DATE: June 13, 1997
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M

REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0074-999
TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEFAX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 276 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-10-092-404-1

Query Match 99.5%; Score 1513; DB 15; Length 276;
Best Local Similarity 99.6%; Pred. No. 8.6e-146;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISSQ 60
DB 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISSQ 60
QY 61 MWLQLSQSLKGDHMTVDFTWMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 61 MWLQLSQSLKGDHMTVDFTWMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
QY 121 QDHLFCPDTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 121 QDHLFCPDTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
QY 181 DQVPLVAVKTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
DB 181 DQVPLVAVKTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
DB 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276

RESULT 3

US-09-981-606-2
Sequence 2, Application US/09981606
Publication No. US20030129595A1
GENERAL INFORMATION:
APPLICANT: Rothenberg et al.
TITLE OF INVENTION: Mutations associated with iron disorders
FILE REFERENCE: 24065-004CON
CURRENT APPLICATION NUMBER: US/09/981,606
CURRENT FILING DATE: 2002-10-16
PRIOR APPLICATION NUMBER: 09/277,457
PRIOR FILING DATE: 1999-03-26
NUMBER OF SEQ ID NOS: 30
SOFTWARE: Patent in Ver. 2.1
SEQ ID NO 2
LENGTH: 348
TYPE: PRT
ORGANISM: Homo sapiens
US-09-981-606-2
Query Match 99.5%; Score 1513; DB 12; Length 348;
Best Local Similarity 99.6%; Pred. No. 1.2e-145;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISSQ 82
QY 61 MWLQLSQSLKGDHMTVDFTWMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGDHMTVDFTWMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142

QY 121 QHLEFCPTDLWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
Db 143 QHLEFCPTDLWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQVPPPLVKVTHVTSSVTLRCALNYYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
Db 203 DQVPPPLVKVTHVTSSVTLRCALNYYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262
QY 241 TYQGWITLAVPPGEORRYTCQVEHPGLDQPLIWIWE 276
Db 263 TYQGWITLAVPPGEORRYTCQVEHPGLDQPLIWIWE 298
RESULT 4
US-10-092-404-3
; Sequence 3, Application US/10092404
; Publication No. US20030073627A1
; GENERAL INFORMATION:
; APPLICANT: Feder, John N.
; Bjorkman, Pamela J.
; Schatzman, Randall C.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
; DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
; AND IRON DEFICIENCY DISEASES
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennire & Edmonds, LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10036-2811
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: Windows
; SOFTWARE: FastSeq for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/10/092,404
; FILING DATE: 04-Mar-2002
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/09/094,964
; FILING DATE: June 12, 1998
; APPLICATION NUMBER: 08/876,010
; FILING DATE: June 13, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0074-999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-493-4935
; TELEFAX: 650-493-5556
; TELEX: 66141 PENNIE
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 276 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; SEQUENCE DESCRIPTION: SEQ ID NO: 3:
US-10-092-404-3

Query Match 98.2%; Score 1493; DB 15; Length 276;
Best Local Similarity 98.9%; Pred. No. 9.3e-144;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVFDDESRRVPRTPWSSRISQ 60
Db 1 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVFDDESRRVPRTPWSSRISQ 60
QY 61 MWLQLSQLKGDHMTFTVDFWTIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDG 120

Db 61 MWLQLSQLKGDHMTFTVDFWTIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDG 120
QY 121 QHLEFCPTDLWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
Db 121 QHLEFCPTDLWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
QY 181 DQVPPPLVKVTHVTSSVTLRCALNYYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
Db 181 DQVPPPLVKVTHVTSSVTLRCALNYYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
QY 241 TYQGWITLAVPPGEORRYTCQVEHPGLDQPLIWIWE 276
Db 241 TYQGWITLAVPPGEORRYTCQVEHPGLDQPLIWIWE 276

RESULT 5
US-10-016-634A-120
; Sequence 120, Application US/10016634A
; Publication No. US20020192666A1
; GENERAL INFORMATION:
; APPLICANT: Sun, Yongming
; APPLICANT: Recipon, Herve
; APPLICANT: Ghosh, Malavika
; APPLICANT: Liu, Chenguang
; TITLE OF INVENTION: Compositions and Methods Relating to Colon Specific Genes and Proteins
; FILE REFERENCE: DEX-0255
; CURRENT APPLICATION NUMBER: US/10/016,634A
; CURRENT FILING DATE: 2001-10-31
; PRIOR APPLICATION NUMBER: US 60/244,258
; PRIOR FILING DATE: 2000-10-31
; NUMBER OF SEQ ID NOS: 176
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 120
; LENGTH: 92
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-016-634A-120

Query Match 33.8%; Score 514; DB 14; Length 92;
Best Local Similarity 100.0%; Pred. No. 9.5e-45;
Matches 92; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY 92 ESHTLQVILGCEMQEDNSTEGYWKYGYDQDHLFCPTDLWRAAEPRAPWTKLEWERHK 151
Db 1 ESHTLQVILGCEMQEDNSTEGYWKYGYDQDHLFCPTDLWRAAEPRAPWTKLEWERHK 60
QY 152 IRARONRAYLERDCPAQLQQLLELGRGVLDQ 183
Db 61 IRARONRAYLERDCPAQLQQLLELGRGVLDQ 92

RESULT 6
US-10-073-300-6
; Sequence 6, Application US/10073300
; Publication No. US20030003535A1
; GENERAL INFORMATION:
; APPLICANT: Reiter, Yoram
; TITLE OF INVENTION: SINGLE CHAIN CLASS I MAJOR HISTO- COMPATIBILITY COMPLEXES
; FILE REFERENCE: 02/23339
; CURRENT APPLICATION NUMBER: US/10/073,300
; CURRENT FILING DATE: 2002-06-25
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 6
; LENGTH: 280
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-073-300-6

Query Match 33.3%; Score 506; DB 15; Length 280;
Best Local Similarity 39.4%; Pred. No. 2.7e-43;
Matches 109; Conservative 45; Mismatches 115; Indels 8; Gaps 7;

APPLICANT: Zhao, Xumei
TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS, KITS, AND
METHODS FOR IDENTIFICATION, ASSESSMENT, PREVENTION, AND
THERAPY OF PROSTATE CANCER
FILE REFERENCE: MRI-044
CURRENT APPLICATION NUMBER: US/10/205,823
CURRENT FILING DATE: 2002-07-25
PRIOR APPLICATION NUMBER: 60/307,982
PRIOR FILING DATE: 2001-07-25
PRIOR APPLICATION NUMBER: 60/314,356
PRIOR FILING DATE: 2001-08-22
PRIOR APPLICATION NUMBER: 60/325,020
PRIOR FILING DATE: 2001-09-25
PRIOR APPLICATION NUMBER: 60/341,746
PRIOR FILING DATE: 2001-12-12
PRIOR APPLICATION NUMBER: 60/362,158
PRIOR FILING DATE: 2002-03-05
NUMBER OF SEQ ID NOS: 455
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 42
LENGTH: 298
TYPE: PRT
ORGANISM: Homo sapiens
US-10-205-823-42

Query Match 32.4%; Score 492; DB 15; Length 298;
Best Local Similarity 36.7%; Pred. No. 7.9e-42;
Matches 101; Conservative 53; Mismatches 111; Indels 10; Gaps 4;
QY 6 HSLHYLFMGASQDGLSLFEALGYDDQLFVYDDSRVPRTPWSSRISQWMLQL 65
DB 28 YSLTYITGLSKHVEDVPAFQALGSLNDLQFFRYNSKDRKSPQMLRWQVE-GMEDWKQD 86
QY 66 SSQLKGDHMTVDFTWIMENHNHSHKESHTLQVILGCENQEDNSTEGYWKYGYDGDHLE 125
DB 87 SLOKAREDFMETLKDIVBYINDSGHVLQGRFCEIENNRSGAFWKYYDGDYIE 146
QY 126 FCPDITLDWRAAPRAWPTKLEWRHKIRARONRAYLERDCPAQLQQLLELGRGLVLDQVVP 185
DB 147 FNKEIPAWVPDPAQITKQWEAPVYVQRAKAYLEECPTALRKLYKSKNILDQDP 206
QY 186 PLVKVT-HVTSVTLRCALNYTPQNTMKWLKDKQPMDAKEPEPK---DVLNPGDGT 241
DB 207 PSVVVTSQAPGKCKLCLAYDFYFGKIDVHWTRAGEVQ-----BPFLRGDVLHNGNGT 261
QY 242 YQGWITLAVPPGGEQRYTCOVHPGLDPLIWIWE 276
DB 262 YQSWVVAVPPQDTAPYSCVHQHSSLAQPLVVPWE 296

RESULT 10
US-10-177-293-23
Sequence 23, Application US/10177293
Publication No. US20030124128A1
GENERAL INFORMATION:
APPLICANT: Liillie, James
APPLICANT: Glatt, Karen
APPLICANT: Zhao, Xumei
APPLICANT: Ganavarapu, Manjula
APPLICANT: Kamatkar, Shubhangi
APPLICANT: Mertens, Maureen
APPLICANT: Myer, Vic
APPLICANT: Wang, Youzhen
APPLICANT: Xu, Yongyao
APPLICANT: Hoersch, Sebastian
APPLICANT: Monahan, John
APPLICANT: Meyers, Rachel E.
APPLICANT: Bast Jr., Robert C.
APPLICANT: Hortobagyi, Gabriel N.
APPLICANT: Pusztai Lajos
APPLICANT: Meric, Funda
APPLICANT: Sahin, Aysegul

APPLICANT: Mills, Gordon B.
TITLE OF INVENTION: COMPOSITIONS, KITS, AND METHODS FOR IDENTIFICATION, ASSESSMENT,
PREVENTION, AND THERAPY OF BREAST CANCER
FILE REFERENCE: MRI-038
CURRENT APPLICATION NUMBER: US/10/177,293
CURRENT FILING DATE: 2002-06-21
PRIOR APPLICATION NUMBER: US 60/299,887
PRIOR FILING DATE: 2001-06-21
PRIOR APPLICATION NUMBER: US 60/301,572
PRIOR FILING DATE: 2001-06-27
PRIOR APPLICATION NUMBER: US 60/306,501
PRIOR FILING DATE: 2001-07-18
PRIOR APPLICATION NUMBER: US 60/325,002
PRIOR FILING DATE: 2001-09-25
PRIOR APPLICATION NUMBER: US 60/362,585
PRIOR FILING DATE: 2002-03-05
PRIOR APPLICATION NUMBER: US 60/xxx,xxx
PRIOR FILING DATE: 2002-05-14
NUMBER OF SEQ ID NOS: 506
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 23
LENGTH: 298
TYPE: PRT
ORGANISM: Homo sapiens
US-10-177-293-23

Query Match 32.4%; Score 492; DB 15; Length 298;
Best Local Similarity 36.7%; Pred. No. 7.9e-42;
Matches 101; Conservative 53; Mismatches 111; Indels 10; Gaps 4;
QY 6 HSLHYLFMGASQDGLSLFEALGYDDQLFVYDDSRVPRTPWSSRISQWMLQL 65
DB 28 YSLTYITGLSKHVEDVPAFQALGSLNDLQFFRYNSKDRKSPQMLRWQVE-GMEDWKQD 86
QY 66 SSQLKGDHMTVDFTWIMENHNHSHKESHTLQVILGCENQEDNSTEGYWKYGYDGDHLE 125
DB 87 SLOKAREDFMETLKDIVBYINDSGHVLQGRFCEIENNRSGAFWKYYDGDYIE 146
QY 126 FCPDITLDWRAAPRAWPTKLEWRHKIRARONRAYLERDCPAQLQQLLELGRGLVLDQVVP 185
DB 147 FNKEIPAWVPDPAQITKQWEAPVYVQRAKAYLEECPTALRKLYKSKNILDQDP 206
QY 186 PLVKVT-HVTSVTLRCALNYTPQNTMKWLKDKQPMDAKEPEPK---DVLNPGDGT 241
DB 207 PSVVVTSQAPGKCKLCLAYDFYFGKIDVHWTRAGEVQ-----EPFLRGDVLHNGNGT 261
QY 242 YQGWITLAVPPGGEQRYTCOVHPGLDPLIWIWE 276
DB 262 YQSWVVAVPPQDTAPYSCVHQHSSLAQPLVVPWE 296

RESULT 11
US-10-015-535-32
Sequence 32, Application US/10015535
Publication No. US20030036506A1
GENERAL INFORMATION:
APPLICANT: Kranz, David M.
APPLICANT: Brophy, Susan
TITLE OF INVENTION: Mutated Class I Major Histocompatibility proteins and
Complexes
FILE REFERENCE: 100-00
CURRENT APPLICATION NUMBER: US/10/015,535
CURRENT FILING DATE: 2001-12-10
PRIOR APPLICATION NUMBER: 60/254,495
PRIOR FILING DATE: 2000-12-08
NUMBER OF SEQ ID NOS: 37
SOFTWARE: Patent in Ver. 2.0
SEQ ID NO 32
LENGTH: 542
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: Synthetic


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; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic
; OTHER INFORMATION: peptide
US-10-015-535-22

Query Match      31.2%; Score 474; DB 15; Length 540;
Best Local Similarity 39.5%; Pred. No. 1.2e-39;
Matches 109; Conservative 40; Mismatches 119; Indels 8; Gaps 7;

QY      6 HSLHYLFMGASEODLGLSFEALGYVDDQLFVYDD--ESRRVEPRTPWSSRISSQMWL 63
Db      144 HSLRYFTAVSRPGLGEPRYMEVGYDDTEFVRPDSDAENPRYEPARMWEQE-GPEYWE 202

QY      64 QLSQSLKGMDFMTVDFTIMENHNHSHK-ESHTLQVILGCEMOEDNS-TEGYWKYGDGQ 121
Db      203 RETQKAKGNEQSRVLDRLTLGGYNQSKGSHTIQVIGCEVSGDGLLRLGYQYAYDGC 262

QY      122 DHLEFCPDTLDWAAPRAWPTKLEWRHKIRARONRAYLERDCPAQLQQLLELGRVLD 181
Db      263 DYIALNEDLKTWTAADMAALITKHKEQAG-EAERLRAYLEGTCEVWLRRLKNGNATLL 321

QY      182 QQVPPPLVKVTHV-TSSVTTLRCALNYYPONITMKWLKDKQPMDAKEFEPKDVLPNGDG 240
Db      322 RTDSPKAVHTHSRPEDKVTLCRCWALGFYPADITLTWQNGEEL-IQDMELVETRPAGDG 380

QY      241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 276
Db      381 TFQKASVVVPLGKEQYITCHVYHQGLPEPLTLRWE 416
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RESULT 15
US-10-015-535-28
; Sequence 28, Application US/10015535
; Publication No. US20030036506A1
; GENERAL INFORMATION:
; APPLICANT: Kranz, David M.
; TITLE OF INVENTION: Mutated Class I Major Histocompatibility proteins and
; FILE REFERENCE: 100-00
; CURRENT APPLICATION NUMBER: US/10/015,535
; PRIOR FILING DATE: 2001-12-10
; PRIOR APPLICATION NUMBER: 60/254,495
; PRIOR FILING DATE: 2000-12-08
; NUMBER OF SEQ ID NOS: 37
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 28
; LENGTH: 541
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic
; OTHER INFORMATION: peptide
US-10-015-535-28

Query Match      31.2%; Score 474; DB 15; Length 541;
Best Local Similarity 39.5%; Pred. No. 1.2e-39;
Matches 109; Conservative 40; Mismatches 119; Indels 8; Gaps 7;

QY      6 HSLHYLFMGASEODLGLSFEALGYVDDQLFVYDD--ESRRVEPRTPWSSRISSQMWL 63
Db      56 HSLRYFTAVSRPGLGEPRYMEVGYDDTEFVRPDSDAENPRYEPARMWEQE-GPEYWE 114

QY      64 QLSQSLKGMDFMTVDFTIMENHNHSHK-ESHTLQVILGCEMOEDNS-TEGYWKYGDGQ 121
Db      115 RETQKAKGNEQSRVLDRLTLGGYNQSKGSHTIQVIGCEVSGDGLLRLGYQYAYDGC 174

QY      122 DHLEFCPDTLDWAAPRAWPTKLEWRHKIRARONRAYLERDCPAQLQQLLELGRVLD 181
Db      175 DYIALNEDLKTWTAADMAALITKHKEQAG-EAERLRAYLEGTCEVWLRRLKNGNATLL 233
```

```
QY      182 QQVPPPLVKVTHV-TSSVTTLRCALNYYPONITMKWLKDKQPMDAKEFEPKDVLPNGDG 240
Db      234 RTDSPKAVHTHSRPEDKVTLCRCWALGFYPADITLTWQNGEEL-IQDMELVETRPAGDG 292

QY      241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 276
Db      293 TFQKASVVVPLGKEQYITCHVYHQGLPEPLTLRWE 328

Search completed: August 5, 2003, 13:21:55
Job time : 34 secs
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GenCore version 5.1.6
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QM protein - protein search, using sw model

Run on: August 5, 2003, 13:06:39 ; Search time 15 Seconds
(without alignments)
1769.504 Million cell updates/sec

Title: US-10-092-404-3
Perfect score: 1514
Sequence: 1 RLRSLSLHFLFMGASEQDL.....RYTCQVEHPGLDPLIVWE 276
Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 283308 seqs, 96168682 residues

Total number of hits satisfying chosen parameters: 283308

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : PIR 76.*

1: pir1.*
2: pir2.*
3: pir3.*
4: pir4.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Match	Length	ID	Description
1	1129	74.6	359	2 JCS382	hereditary hemochromatosis precursor - mouse
2	530.5	35.0	341	2 A57136	class I histocomp
3	517	34.1	361	1 HLRB	MHC class I histoc
4	517	34.1	361	2 I46858	MHC class I histoc
5	514	33.9	332	2 S06424	MHC class I histoc
6	511	33.8	365	2 I36961	MHC class I histoc
7	510	33.7	361	2 B27638	MHC class I histoc
8	509	33.6	365	2 I83063	HLA-A*0205 pr
9	508	33.6	365	2 A47636	MHC class I histoc
10	508	33.6	365	2 I56039	HLA-A*03.3 precursor
11	506	33.4	370	1 HLHUA3	MHC class I histoc
12	504	33.3	365	2 I38439	MHC class I histoc
13	503	33.2	365	2 I37542	MHC class I histoc
14	503	33.2	365	2 I38442	gene HLA-A*0205 pr
15	503	33.2	365	2 I61902	MHC class I histoc
16	502	33.2	365	2 I72170	MHC class I histoc
17	502	33.2	365	2 I38441	gene HLA-A*6802 pr
18	501	33.1	355	2 T28149	MHC class I histoc
19	500	33.0	365	1 HLHUA2	MHC class I histoc
20	500	33.0	365	2 I37482	MHC class I histoc
21	500	33.0	365	2 I38519	MHC class I histoc
22	500	33.0	365	2 I84448	MHC class I histoc
23	499	33.0	365	2 I38610	MHC class I histoc
24	499	33.0	365	2 I37470	HLA-A*0210 - human
25	498	32.9	364	2 S03535	MHC class I histoc
26	497	32.8	365	2 I37476	MHC class I histoc
27	497	32.8	365	2 I37478	MHC class I histoc
28	497	32.8	365	2 I38443	Gene HLA-A*0203 pr
29	497	32.8	365	2 I61857	MHC HLA-A2.4a chain

MHC class I protei
MHC class I histoc
MHC class I histoc
MHC class I histoc
MHC class I histoc
MHC class I histoc
zinc-alpha 2-glyco
MHC class I histoc
class I histocompa
HLA-AW34.2 antigen
HLA-AW34.2 antigen
MHC class I histoc
MHC class I histoc
HLA-AW33.1, HLA-AW
MHC class I histoc
MHC class I histoc
major histocompati

ALIGNMENTS

RESULT 1

JCS382 hereditary hemochromatosis protein precursor - mouse

C;Species: Mus musculus (house mouse)

C;Date: 02-Jun-1997 #sequence_revision 18-Jul-1997 #text_change 05-Nov-1999

C;Accession: JCS382

R;Hashimoto, K.; Hirai, M.; Kurosawa, Y.

Biochem. Biophys. Res. Commun. 230, 35-39, 1997

A;Title: Identification of a mouse homolog for the human hereditary haemochromatosis ca

A;Reference number: JCS382; MUID:97148566; PMID:9020055

A;Accession: JCS382

A;Status: nucleic acid sequence not shown

A;Molecule type: DNA

A;Residues: 1-359 <HAS>

A;Cross-references: GB:U68849; NID:gl519484; PIDN:AAB07525.1; PID:gl519485

C;Comment: This protein plays a role in iron metabolism.

C;Genetics:

A;Gene: mr2

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

F;1-29/Domain: signal sequence #status predicted <SIG>

F;30-359/Product: hereditary haemochromatosis protein #status predicted <MAT>

F;30-117/Domain: alpha 1 #status predicted <ALF1>

F;118-217/Domain: alpha 2 #status predicted <ALF2>

F;218-309/Domain: alpha 3 #status predicted <ALF3>

F;314-340/Domain: transmembrane #status predicted <TRM>

F;341-359/Domain: intracellular #status predicted <INT>

Query Match 74.6%; Score 1129; DB 2; Length 359;

Best Local Similarity 71.9%; Pred. No. 1.9e-86;

Matches 202; Conservative 30; Mismatches 41; Indels 8; Gaps 1;

QY 4 RSHSLHFLFMGASEQDLGLSLFEALGYVDDQLFVYDHSRVRVPTWVSRISSQWL 63

DB 30 RSHSLHFLFMGASEQDLGLSLFEALGYVDDQLFVYDHSRVRVPTWVSRISSQWL 89

QY 64 QLSQSLKGDHMTFTVDFTIMENHASK-----ESHTLQVLGCMEQDNSTEGYWK 115

DB 90 HLSQSLKGDHMTFTVDFTIMENHASK-----ESHTLQVLGCMEQDNSTEGYWK 149

QY 116 YGYDQDLEFCPTDLDRRAEPRAWPTKLEWERHKIRARQNYLERDCAQLOQLLEL 175

DB 150 YGYDQDLEFCPTDLDRRAEPRAWPTKLEWERHKIRARQNYLERDCAQLOQLLEL 209

QY 176 GSGVLDQVPLAVKTHVTSSVTTLRCALNYYPQNTMKWLKQKPMDAKEPEPKDVL 235

DB 210 GSGVLDQVPLAVKTHVTSSVTTLRCALNYYPQNTMKWLKQKPMDAKEPEPKDVL 269

QY 236 PNGDGTQGWITLAVPFGEEQRYTCQVEHPGLDQPLIVWE 276

DB 270 PNGDGTQGWITLAVPFGEEQRYTCQVEHPGLDQPLIVWE 310


```

F;110/Binding site: carbohydrate (Asn) (covalent) #status predicted
F;125-188,227-283/Disulfide bonds: #status predicted

Query Match          34.1%; Score 517; DB 1; Length 361;
Best Local Similarity 40.1%; Pred. No. 1.7e-35;
Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7

QY 5 SHSLHYLFMGASQDGLSLFEALGYVDDQLGFVYDHE--SRRVEPTPTWVSSRISSQW 62
   |||: : : : : |||: : : : : |||: : : : : |||: : : : : |||: : : : :
Db 26 SHSMRYFYTSVRPGLGEPRFIIVGYDDTQFVRFDSDAASPRMEQAPWM-GQVEPEYW 84

QY 63 LQLSOSLKGWDHMTVDFTWIMENHNASKB-SHTLQVILGCEMOEDNS-TGYWKYGYDG 120
   |||: : : : : |||: : : : : |||: : : : : |||: : : : : |||: : : : :
Db 85 DQQTQIAKTAQTAQFRVNLNTALRYNQSAAGSHTFQTMFGCEVWADGRFFHYGKYQAYDG 144

QY 121 QDALEFCPTDLWDRAAEPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
   |||: : : : : |||: : : : : |||: : : : : |||: : : : : |||: : : : :
Db 145 ADYIALNEDLRSTWADTAQNTQKWEAAG-EAERHRAVLERECVWLRYLEMGKETL 203

QY 181 DQGVPLPVKVVTHVTS-VTTLCRALNYYPNITMKWLKDKQPMDAKEPFEPKDVLPNGD 239
   |||: : : : : |||: : : : : |||: : : : : |||: : : : : |||: : : : :
Db 204 QRADPPKAHVTHHPASDREATLRCWALGFYPAISLTWQDGED-QTQDTLAVETRPQGD 262

QY 240 GTYQGHITLAVPGEQRVTCVHEPGLDQPLVIWE 276
   |||: : : : : |||: : : : : |||: : : : : |||: : : : : |||: : : : :
Db 263 GTFQKAAVVPVGEQRVTCRVOHEGLPEPLTLTWE 299

RESULT 4
I46858
MHC class I RIA precursor - rabbit
C:Species: Oryctolagus cuniculus (domestic rabbit)
C:Date: 14-Feb-1997 #sequence_revision 14-Feb-1997 #text_change 21-Jan-2000
C:Accession: I46858
R:Marche, P.N.; Tykocinski, M.L.; Max, E.E.; Kindt, T.J.
Immunogenetics 21, 71-82, 1985
A:Title: Structure of a functional rabbit class I MHC gene: Similarity to human
A:Reference number: I46858; MUID:85103547; PMID:3917974
A:Accession: I46858
A:Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: DNA
A:Residues: 1-361 <VAR>
A:Cross-references: GB:K02819; NID:g165497; PIDN:AAA98730.1; PID:g165498
C:Genetics:
A:Introns: 25/1; 115/1; 207/1; 295/1; 337/1; 348/1
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
F;220-285/Domain: immunoglobulin homology <IMM>

Query Match          34.1%; Score 517; DB 2; Length 361;
Best Local Similarity 40.1%; Pred. No. 1.7e-35;
Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7

QY 5 SHSLHYLFMGASQDGLSLFEALGYVDDQLGFVYDHE--SRRVEPTPTWVSSRISSQW 62
   |||: : : : : |||: : : : : |||: : : : : |||: : : : : |||: : : : :
Db 26 SHSMRYFYTSVRPGLGEPRFIIVGYDDTQFVRFDSDAASPRMEQAPWM-GQVEPEYW 84

QY 63 LQLSOSLKGWDHMTVDFTWIMENHNASKB-SHTLQVILGCEMOEDNS-TGYWKYGYDG 120
   |||: : : : : |||: : : : : |||: : : : : |||: : : : : |||: : : : :
Db 85 DQQTQIAKTAQTAQFRVNLNTALRYNQSAAGSHTFQTMFGCEVWADGRFFHYGKYQAYDG 144

QY 121 QDALEFCPTDLWDRAAEPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
   |||: : : : : |||: : : : : |||: : : : : |||: : : : : |||: : : : :
Db 145 ADYIALNEDLRSTWADTAQNTQKWEAAG-EAERHRAVLERECVWLRYLEMGKETL 203

QY 181 DQGVPLPVKVVTHVTS-VTTLCRALNYYPNITMKWLKDKQPMDAKEPFEPKDVLPNGD 239
   |||: : : : : |||: : : : : |||: : : : : |||: : : : : |||: : : : :
Db 204 QRADPPKAHVTHHPASDREATLRCWALGFYPAISLTWQDGED-QTQDTLAVETRPQGD 262

QY 240 GTYQGHITLAVPGEQRVTCVHEPGLDQPLVIWE 276
   |||: : : : : |||: : : : : |||: : : : : |||: : : : : |||: : : : :
Db 263 GTFQKAAVVPVGEQRVTCRVOHEGLPEPLTLTWE 299

```

RESULT 5

S06424
MHC class I histocompatibility antigen Ch25 alpha chain precursor - chimpanzee
N;Alternate names: MHC Ch1A chain
C;Species: Pan troglodytes (chimpanzee)
C;Date: 19-Mar-1997 #sequence_revision 19-Mar-1997 #text_change 23-Jul-1999
C;Accession: S06424; I36959
R;Lawlor, D.A.; Ward, F.E.; Ennis, P.D.; Jackson, A.P.; Parham, P.
Nature 335, 268-271, 1988
A;Title: HLA-A and B polymorphisms predate the divergence of humans and chimpanzees.
A;Reference number: S06424; MUID:98319000; PMID:3412487
A;Accession: S06424
A;Molecule type: mRNA
A;Residues: 1-332 <LAW>
R;Parham, P.; Lawlor, D.A.; Lomen, C.E.; Ennis, P.D.
J. Immunol. 142, 3937-3950, 1989
A;Title: Diversity and diversification of HLA-A,B,C alleles.
A;Reference number: I36956; MUID:99235215; PMID:2715640
A;Accession: I36959
A;Molecule type: mRNA
A;Residues: 1-332 <RES>
A;Cross-references: GB:M24047; NID:g176818; PID:AAA35426.1; PID:g553155
C;Superfamily: Class I histocompatibility antigen; immunoglobulin homology
C;Keywords: glycoprotein; membrane protein
F;1-24/Domain: signal sequence #status predicted <SIG>
F;25-114/Domain: alpha-1 #status predicted
F;115-206/Domain: alpha-2 #status predicted <EX1>
F;220-285/Domain: immunoglobulin homology <IMM>
F;307-331/Domain: transmembrane #status predicted <TM>
F;110/Binding site: carbohydrate (Asn) (covalent) #status predicted
F;125-188, 227-283/Disulfide bonds: #status predicted

Query Match 33.9%; Score 514; DB 2; Length 332;
Best Local Similarity 40.1%; Pred. No. 2.7e-35;
Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7;

QY 5 SLSLHLYFMGASQDGLSLFEALGVDDQLFVFDHE--SRVPRTPWSSRISQMW 62

DB 26 SHSMRYFFTSVSRPGRGEPFIAVGVDVDTQFVRFDSDAASQRMPEAPWIEQ-GPEYW 84

QY 63 LQLSQLKGDHMTVDFTWIMENHNASKE-SHTLQVILGCEMOEDNS-TEGYWKYGDG 120

DB 85 DQTRAKAHSQTRVDLGLTRGYNQSDGSHTIQIMYGCDDVGSQGRFLRGTRQDAYDG 144

QY 121 QDALEFCPTDLWRAAPRAWPTKLEWERHKIRARQNRAYLIERDCPAQLQQLLELGRGVL 180

DB 145 KDVIALLNEDLRSWTAADMAAQITKRWAAH-AAEQRAYLEGTCVLEWLRRLYENGKEYL 203

QY 181 DQOVPLVKTTH-VTSSVTTLRCALNYYPQNIWKMLKDKQPMDAKFEPEKDVLPNGD 239

DB 204 QRTDPPKTHHTHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTLVELVETRPAGD 262

QY 240 GTYQGWITLAVPGEQRYTCQVEHPGLDPLIVWE 276

DB 263 GTFOKAAVVPVSGEQRVTCHVQHEGLPKPLTLRWE 299

RESULT 6

I36961

MHC class I protein - chimpanzee

C;Species: Pan troglodytes (chimpanzee)

C;Date: 04-Oct-1996 #sequence_revision 04-Oct-1996 #text_change 21-Jan-2000

C;Accession: I36961

R;Lawlor, D.A.; Warren, E.; Ward, F.E.; Parham, P.

Immunol. Rev. 113, 147-185, 1990

A;Title: Comparison of class I MHC alleles in humans and apes.

A;Reference number: I36961; MUID:90201944; PMID:1690682

A;Accession: I36961

A;Status: preliminary; translated from GB/EMBL/DDBJ

A;Molecule type: mRNA

A;Residues: 1-365 <RES>

A;Cross-references: GB:M30678; NID:g176822; PID:AAA87970.1; PID:g176823

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

Query Match 33.9%; Score 514; DB 2; Length 332;
Best Local Similarity 40.1%; Pred. No. 2.7e-35;
Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7;

QY 5 SLSLHLYFMGASQDGLSLFEALGVDDQLFVFDHE--SRVPRTPWSSRISQMW 62

DB 26 SHSMRYFFTSVSRPGRGEPFIAVGVDVDTQFVRFDSDAASQRMPEAPWIEQ-GPEYW 84

QY 63 LQLSQLKGDHMTVDFTWIMENHNASKE-SHTLQVILGCEMOEDNS-TEGYWKYGDG 120

DB 85 DQTRAKAHSQTRVDLGLTRGYNQSDGSHTIQIMYGCDDVGSQGRFLRGTRQDAYDG 144

QY 121 QDALEFCPTDLWRAAPRAWPTKLEWERHKIRARQNRAYLIERDCPAQLQQLLELGRGVL 180

DB 145 KDVIALLNEDLRSWTAADMAAQITKRWAAH-AAEQRAYLEGTCVLEWLRRLYENGKEYL 203

QY 181 DQOVPLVKTTH-VTSSVTTLRCALNYYPQNIWKMLKDKQPMDAKFEPEKDVLPNGD 239

DB 204 QRTDPPKTHHTHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTLVELVETRPAGD 262

QY 240 GTYQGWITLAVPGEQRYTCQVEHPGLDPLIVWE 276

DB 263 GTFOKAAVVPVSGEQRVTCHVQHEGLPKPLTLRWE 299

RESULT 7

B27638

MHC class I histocompatibility antigen alpha chain precursor (BL3-7) - bovine

C;Species: Bos primigenius taurus (cattle)

C;Date: 08-Mar-1989 #sequence_revision 08-Mar-1989 #text_change 16-Feb-1997

C;Accession: B27638

R;Ennis, P.D.; Jackson, A.P.; Parham, P.

J. Immunol. 141, 642-651, 1988

A;Title: Molecular cloning of bovine class I MHC cDNA.

A;Reference number: A92826; MUID:89258075; PMID:3133413

A;Accession: B27638

A;Status: not compared with conceptual translation

A;Molecule type: mRNA

A;Residues: 1-361 <ENN>

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

C;Keywords: heterodimer; transmembrane protein

F;1-24/Domain: signal sequence #status predicted <SIG>

F;25-361/Product: MHC class I histocompatibility antigen, BoLA alpha chain (BL3-7) #stat

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.7%; Score 510; DB 2; Length 361;
Best Local Similarity 38.9%; Pred. No. 6.4e-35;
Matches 109; Conservative 49; Mismatches 114; Indels 8; Gaps 7;

QY 2 LRLSHLYLFMGASEQDGLSLFEALGVDDQLFVFDHE--SRVPRTPWSSRIS 59

DB 23 LAGSHLYFTYGVSRPGLGEPFIAVGVDVDTQFVRFDSDAPNPREBVRVPMSE-QP 81

QY 60 QMWLQSLKGDHMTVDFTWIMENHNASKE-SHTLQVILGCEMOEDNS-TEGYWKY 117

DB 82 EYWDNTRYIKDTAQIFRVDLNLRGYNNQSGTSHNIQAMYGVCDVGDGRLLRGFQFG 141

QY 118 YDQDALEFCPTDLWRAAPRAWPTKLEWERHKIRARQNRAYLIERDCPAQLQQLLELGR 177

DB 142 YDGRDYIALNEELRSWTAADTAQAQITKRWAAH-AAETWRNLYSGECVLEWLRRLYENG 200

QY 178 GVLDQOVPLVKTTH-VTSSVTTLRCALNYYPQNIWKMLKDKQPMDAKFEPEKDVLP 236

DB 201 DTLRADPPKHAHVTHHSIDREVTILRCWALGFYPEEISLTWQREGED-QTQDMELVETRP 259

QY 237 NGDGYTQGWITLAVPGEQRYTCQVEHPGLDPLIVWE 276

DB 260 SGDGTFOKAAVVPVSGEQRVTCHVQHEGLPKPLTLRWE 299

RESULT 8

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.8%; Score 511; DB 2; Length 365;
Best Local Similarity 39.7%; Pred. No. 5.4e-35;
Matches 110; Conservative 44; Mismatches 115; Indels 8; Gaps 7;

QY 5 SLSLHLYFMGASEQDGLSLFEALGVDDQLFVFDHE--SRVPRTPWSSRISQMW 62

183063
 All.1.2 - human
 C:Species: Homo sapiens (man)
 C>Date: 02-Aug-1996 #sequence_revision 02-Aug-1996 #text_change 21-Jan-2000
 C:Accession: I83063
 R:Lin, L.; Tokunaga, K.; Ishikawa, Y.; Bannai, M.; Kashiwase, K.; Kuwata, S.; Akaza, T.;
 Tissue Antigens 43, 78-82, 1994
 A:Title: Sequence analysis of serological HLA-A11 split antigens, All.1.1 and All.1.2.
 A:Reference number: I60129; MUID:94287401; PMID:8016845
 A:Accession: I83063
 A>Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: mRNA
 A:Residues: 1-365 <RES>
 A:Cross-references: GB:D16842; NID:G540517; PIDN:BA04118.1; PID:G487911
 C:Genetics:
 A:Gene: A1102
 C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
 F:220-285/Domain: immunoglobulin homology <IMM>
 Query Match 33.6%; Score 509; DB 2; Length 365;
 Best Local Similarity 39.4%; Pred. No. 7.9e-35;
 Matches 109; Conservative 46; Mismatches 114; Indels 8; Gaps 7;
 QY 5 SHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDHE--SRVPRTPWVSSRISSQW 62
 DB 26 SHSMRYFYTSVRPGKPRFIAVGYVDDTFVRFSDAASQRMPEAPWIEQE-GPEYW 84
 QY 63 LQLSQSLKGDHMFVDFWTIMENASKE-SHTLQVILGCMEQDNS-TEGWYKYGDG 120
 DB 85 DQSTRNVKAQSDTRVDLGLTRGYNQSDGSHTIQIMYCGDVPGRFLRGYDAYDG 144
 QY 121 QDALEFCPDLDRWAAPRAWPYKLEWERHKIRARONRAYLERDCPAQLQQLLELGRV 180
 DB 145 KDVALNEDLRSTWADMAAQITKRWEAAH-AEQRAYLEGRCVWELRYLNGKETL 203
 QY 181 DQVPPPLVKVTHH-VTSSVTLRCALNYYPNITMKWKDKQPMDAKEPEPKDVLNPGD 239
 DB 204 QRTDPPKTHMTHPISDHEATLRCWALGYFPAEITLTWQDGED-QTQDTVELVETRPAGD 262
 QY 240 GTYQGWITLAVPGEORVTCVHEPGLDQPLIVWE 276
 DB 263 GTFOKAAVVPVSGEORVTCVHQHEGLPKPLTRWE 299
 RESULT 9
 A47636
 MHC class I histocompatibility antigen HLA-A11 alpha chain precursor - human
 C:Species: Homo sapiens (man)
 C>Date: 31-Dec-1993 #sequence_revision 28-Apr-1995 #text_change 23-Jul-1999
 C:Accession: S03536; S03694; A47636; I60129
 R:Mayer, W.E.; Jonker, M.; Klein, D.; Ivanyi, P.; van Seventer, G.; Klein, J.
 EMO J. 7, 2765-2774, 1988
 A:Title: Nucleotide sequences of chimpanzee MHC class I alleles: evidence for trans-spec
 A:Reference number: S01171; MUID:89030641; PMID:2460344
 A:Accession: S03536
 A:Molecule type: mRNA
 A:Residues: 1-365 <WAY>
 A:Cross-references: EMBL:X13111; NID:G32138; PIDN:CAA31503.1; PID:G32139
 A>Note: this allele is designated A*1101 (formerly A11E, A11.1)
 A:Accession: S03694
 A:Molecule type: mRNA
 A:Residues: 1-42, 'K', 44-298 <WA2>
 A:Cross-references: EMBL:X13112; NID:G32142; PIDN:CAA31504.1; PID:G32143
 A>Note: this allele is designated A*1102 (formerly A11K, A11.2)
 R:Cowan, E.P.; Jelachich, M.L.; Biddison, W.E.; Coligan, J.E.
 Immunogenetics 25, 241-250, 1987
 A:Title: DNA sequence of HLA-A11: remarkable homology with HLA-A3 allows identification
 A:Reference number: A47636; MUID:87192928; PMID:2437024
 A:Accession: A47636
 A:Molecule type: DNA
 A:Residues: 26-365 <COW>
 A:Cross-references: GB:M16007; GB:M16008; GB:M16009; GB:M16010; NID:G184130; PIDN:AAA654
 A>Note: the authors translated the codon GAC for residue 89 as Ala, CCG for residue 104

A>Note: this allele is designated A*1101 (formerly A11E, A11.1)
 R:Lin, L.; Tokunaga, K.; Ishikawa, Y.; Bannai, M.; Kashiwase, K.; Kuwata, S.; Akaza, T.;
 Tissue Antigens 43, 78-82, 1994
 A:Title: Sequence analysis of serological HLA-A11 split antigens, All.1.1 and All.1.2.
 A:Reference number: I60129; MUID:94287401; PMID:8016845
 A:Accession: I60129
 A>Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: mRNA
 A:Residues: 1-365 <RES>
 A:Cross-references: GB:D16841; NID:G540516; PIDN:BA04117.1; PID:G487909
 A>Note: this allele is designated A*1101 (formerly A11E, A11.1)
 C:Genetics:
 A:Gene: GDB:HLA-A
 A:Cross-references: GDB:119310; OMIM:142800
 A:Map position: 6p21.3-6p21.3
 C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
 C:Keywords: transmembrane protein
 F:1-24/Domain: signal sequence #status predicted <SIG>
 F:25-365/Product: class I histocompatibility antigen alpha chain #status predicted <EXT>
 F:25-298/Domain: extracellular #status predicted <EXT>
 F:220-285/Domain: immunoglobulin homology <IMM>
 F:299-337/Domain: transmembrane #status predicted <TMM>
 F:338-365/Domain: intracellular #status predicted <INT>
 Query Match 33.6%; Score 508; DB 2; Length 365;
 Best Local Similarity 39.4%; Pred. No. 9.5e-35;
 Matches 109; Conservative 46; Mismatches 114; Indels 8; Gaps 7;
 QY 5 SHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDHE--SRVPRTPWVSSRISSQW 62
 DB 26 SHSMRYFYTSVRPGKPRFIAVGYVDDTFVRFSDAASQRMPEAPWIEQE-GPEYW 84
 QY 63 LQLSQSLKGDHMFVDFWTIMENASKE-SHTLQVILGCMEQDNS-TEGWYKYGDG 120
 DB 85 DQSTRNVKAQSDTRVDLGLTRGYNQSDGSHTIQIMYCGDVPGRFLRGYDAYDG 144
 QY 121 QDALEFCPDLDRWAAPRAWPYKLEWERHKIRARONRAYLERDCPAQLQQLLELGRV 180
 DB 145 KDVALNEDLRSTWADMAAQITKRWEAAH-AEQRAYLEGRCVWELRYLNGKETL 203
 QY 181 DQVPPPLVKVTHH-VTSSVTLRCALNYYPNITMKWKDKQPMDAKEPEPKDVLNPGD 239
 DB 204 QRTDPPKTHMTHPISDHEATLRCWALGYFPAEITLTWQDGED-QTQDTVELVETRPAGD 262
 QY 240 GTYQGWITLAVPGEORVTCVHEPGLDQPLIVWE 276
 DB 263 GTFOKAAVVPVSGEORVTCVHQHEGLPKPLTRWE 299
 RESULT 10
 I56039
 HLA-A30.3 precursor - human
 C:Species: Homo sapiens (man)
 C>Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 21-Jan-2000
 C:Accession: I56039
 R:Kato, K.; Trapani, J.A.; Allopenna, J.; Dupont, B.; Yang, S.Y.
 J. Immunol. 143, 3371-3378, 1989
 A:Title: Molecular analysis of the serologically defined HLA-Aw19 antigens. A genet
 A:Reference number: I56039; MUID:90038496; PMID:2478623
 A:Accession: I56039
 A>Status: preliminary; translated from GB/EMBL/DBJ
 A:Molecule type: DNA
 A:Residues: 1-365 <RES>
 A:Cross-references: GB:M30576; NID:G187646; PIDN:AAA59612.1; PID:G386878
 C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
 F:220-285/Domain: immunoglobulin homology <IMM>
 Query Match 33.6%; Score 508; DB 2; Length 365;
 Best Local Similarity 39.4%; Pred. No. 9.5e-35;
 Matches 109; Conservative 47; Mismatches 113; Indels 8; Gaps 7;
 QY 5 SHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDHE--SRVPRTPWVSSRISSQW 62

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Db 26 SHSMRYFFTSVRPGSGEPRIAGVYDDTQFVRFDSDAASQRMPEPRAPWIEQ-EPEYW 84
QY 63 LQLSLSKLGNDHMTFTVDFTWIMENHNASK-SHTLQVILGCEMQEDNS--TEGYWKYGYDG 120
Db 85 DQETRNVAQSQDTRDVLGTLRGYNSQAGSHTIQIMYGCDVSGDGRFLRGYEQHAYDG 144
QY 121 QDALEFCPTDLDWRAAEPRAPWPTKLEWERHUKIRARQNRAVLERDPCPAQLQQLLELGRGVL 180
Db 145 KDYIALNEDLRSWTAADMAAQITQRKWEAAR-WAEQLRAYLEGTCVWELRYLENGKETL 203
QY 181 DQOVPPPLVKVTHH-VTSSVTLRCALNYPONITMKWLKDQKPMDAKEFEPKDVLPNGD 239
Db 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQRDGED-QTQDTLVELTRPADG 262
QY 240 GTYQGWITLAVPPGGEQRYTCQVEHPGLDQPLIVWE 276
Db 263 GTFQKAAVAVVPSGGEQRYTCHVQHEGLPKPLTLRWE 299

RESULT 11
HLHUA3
MHC class I histocompatibility antigen HLA-A3 alpha chain precursor - human
C:Species: Homo sapiens (man)
C>Date: 17-Mar-1987 #sequence_revision 17-Mar-1987 #text_change 02-Sep-1997
C:Accession: A02192
R:Strachan, T.; Sodoyer, R.; Damotte, M.; Jordan, B.R.
EMBO J. 3, 887-894, 1984
A:Title: Complete nucleotide sequence of a functional class I HLA gene, HLA-A3: implicated
A:Reference number: A02192; MUID:84207948; PMID:6609814
A:Accession: A02192
A:Molecule type: DNA
A:Residues: 1-370 <STR>
C:Genetics:
A:Gene: GDB:HLA-A
A:Cross-references: GDB:119310; OMIM:142800
A:Map position: 6p21.3-6p21.3
A:Introns: 30/1; 120/1; 212/1; 304/1; 343/1; 354/1; 370/1
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
C:Keywords: duplication; glycoprotein; heterodimer; transmembrane protein; transplantati
F;1-29/Domain: signal sequence #status predicted <SIG>
F;30-370/Product: class I histocompatibility antigen HLA-A3 alpha chain #status predicted
F;30-312/Domain: extracellular #status predicted <EX1>
F;30-119/Domain: alpha-1 <EX2>
F;120-211/Domain: alpha-2 <EX2>
F;225-290/Domain: immunoglobulin homology <IMM>
F;313-337/Domain: transmembrane #status predicted <TM>
F;338-370/Domain: intracellular #status predicted <INT>
F;115/Binding site: carbohydrate (Asn) (covalent) #status predicted
F;232-288/Disulfide bonds: #status predicted

Query Match 33.4%; Score 506; DB 1; Length 370;
Best Local Similarity 39.6%; Pred. No. 1.4e-34;
Matches 110; Conservative 46; Mismatches 112; Indels 10; Gaps 8;

QY 5 SHSLHYLFMGASQDGLSLFEALGVYDDQLFVFDHE--SRVPRTPWVSSRISSQW 62
Db 31 SHSMRYFFTSVRPGSGEPRIAGVYDDTQFVRFDSDAASQRMPEPRAPWIEQ-EPEYW 89
QY 63 LQLSLSKLGNDHMTFTVDFTWIMENHNASK-SHTLQVILGCEMQEDNS--TEGYWKYGYDG 120
Db 90 DQETRNVAQSQDTRDVLGTLRGYNSQAGSHTIQIMYGCDVSGDGRFLRGYEQHAYDG 149
QY 121 QDALEFCPTDLDWRAAEPRAPWPTKLEWERHUKIRARQNRAVLERDPCPAQLQQLLELGRGV 179
Db 150 KDYIALNEDLRSWTAADMAAQITQRKWEAAR-WAEQLRAYLEGTCVWELRYLENGKET 207
QY 180 LDOQVPPPLVKVTHH-VTSSVTLRCALNYPONITMKWLKDQKPMDAKEFEPKDVLPNG 238
Db 208 LQRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQRDGED-QTQDTLVELTRPADG 266
QY 239 DGTGQWITLAVPPGGEQRYTCQVEHPGLDQPLIVWE 276
Db 267 DGTGQKAAVAVVPSGGEQRYTCHVQHEGLPKPLTLRWE 304
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RESULT 12

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I38439
MHC class I histocompatibility antigen HLA-A*8001 precursor - human
C:Species: Homo sapiens (man)
C>Date: 07-Jun-1996 #sequence_revision 07-Jun-1996 #text_change 21-Jan-2000
C:Accession: I59638; I38439
R:Domena, J.D.; Hildebrand, W.H.; Bias, W.B.; Parham, P.
Tissue Antigens 42, 156-159, 1993
A:Title: A sixth family of HLA-A alleles defined by HLA-A*8001.
A:Reference number: I59638; MUID:94112691; PMID:8284791
A:Accession: I59638
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-365 <DOM>
A:Cross-references: GB:118898; NID:G306853; PIDN:AAAL7012.1; PID:G306854
R:Balas, A.; Garcia-Sanchez, F.; Gomez-Reino, F.; Vicario, J.L.
Immunogenetics 39, 452, 1994
A:Title: Characterization of a new and highly distinguishable HLA-A allele in a Spanish
A:Reference number: I38439; MUID:94245293; PMID:8188325
A:Accession: I38439
A>Status: preliminary; translated from GB/EMBL/DBJ
A:Molecule type: mRNA
A:Residues: 1-365 <BAL>
A:Cross-references: EMBL:U03754; NID:G432407; PIDN:AAOC4322.1; PID:G432408
C:Genetics:
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A:Gene: GDB:HLA-A
A:Cross-references: GDB:119310; OMIM:142800
A:Map position: 6p21.3-6p21.3
C:Superfamily: class I histocompatibility antigen; immunoglobulin homology
F;220-285/Domain: immunoglobulin homology <IMM>
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Query Match 33.3%; Score 504; DB 2; Length 365;

Best Local Similarity 38.3%; Pred. No. 2.1e-34;

Matches 106; Conservative 52; Mismatches 111; Indels 7;

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QY 5 SHSLHYLFMGASQDGLSLFEALGVYDDQLFVFDHE--SRVPRTPWVSSRISSQW 62
Db 26 SHSMRYFFTSVRPGSGEPRIAGVYDDTQFVRFDSDAASQRMPEPRAPWIEQ-EPEYW 84
```

```
QY 63 LQLSLSKLGNDHMTFTVDFTWIMENHNASK-SHTLQVILGCEMQEDNS--TEGYWKYGYDG 120
Db 85 DEETRNVAQSQDTRDVLGTLRGYNSQAGSHTIQIMYGCDVSGDGRFLRGYEQHAYDG 144
```

```
QY 121 QDALEFCPTDLDWRAAEPRAPWPTKLEWERHUKIRARQNRAVLERDPCPAQLQQLLELGRGVL 180
Db 145 KDYIALNEDLRSWTAADMAAQITQRKWEAAR-WAEQLRAYLEGTCVWELRYLENGKETL 203
```

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QY 181 DQOVPPPLVKVTHH-VTSSVTLRCALNYPONITMKWLKDQKPMDAKEFEPKDVLPNGD 239
Db 204 QRTDPPKTHMTHHPISDHEATLRCWALGFYPAEITLTWQRDGED-QTQDTLVELTRPADG 262
```

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QY 240 GTYQGWITLAVPPGGEQRYTCQVEHPGLDQPLIVWE 276
Db 263 GTFQKAAVAVVPSGGEQRYTCHVQHEGLPKPLTLRWE 299
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RESULT 13

I37542

MHC class I histocompatibility antigen HLA-A2 alpha chain (allele A*0216) precursor - h

C:Species: Homo sapiens (man)

C>Date: 04-Oct-1996 #sequence_revision 04-Oct-1996 #text_change 21-Jan-2000

C:Accession: I37542; 849582

R:Barouch, D.; Krausa, P.; Bodmer, J.; Browning, M.J.; McMichael, A.J.

Immunogenetics 41, 388, 1995

A:Title: Identification of a novel HLA-A2 subtype, HLA-A*0216.

A:Reference number: I37542; MUID:95278976; PMID:7759139

A:Accession: I37542

A>Status: preliminary; translated from GB/EMBL/DBJ

A:Molecule type: mRNA

A:Residues: 1-365 <RES>

A:Cross-references: EMBL:Z46633; NID:G575248; PIDN:CAA86602.1; PID:G575249

A;Note: submitted to the EMBL Data Library, November 1994

C;Gene: hla-A

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.2%; Score 503; DB 2; Length 365;
Best Local Similarity 39.4%; Pred. No. 2.5e-34;
Matches 109; Conservative 45; Mismatches 115; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASQDGLSLFEALGVYDDQLFVFDHE--SRVPRTPWSSRISSQMW 62

Db 26 SHSMRYFTTSVRGEPGFIAVGVDQTFVRFDSDAASQRMPEAPWIEQE-GPEYW 84

QY 63 LQLSLSLKGWDHMTFTVDFTIMENHNASKB-SHTLQVILGCMEQED-NSTEGYWKYGYDG 120

Db 85 DGETRKKVKAHSQTHRVDLGLTRGYNQSEAGSHTVQRMVGCVDGSDWRFLRGVHQYAYDG 144

QY 121 QDALEFCPDTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180

Db 145 KDYIALKEDLRSWTAADMAAQTTHKWEAAHV-AEQRLAYLEGCEVWLRRLRYLENGKETL 203

QY 181 DQOVPLPVKVTTH-VTSSVTLTLCRALNYPQNTMKWLKDKQPMDAKFEPEKDVLPNGD 239

Db 204 QRTDAPKTHMTHAVSDHEATLRCWALSFPABITLTWQDGDG-OTQDTLVETRPAGD 262

QY 240 GTYQGHITLAVPGEQRVTCVHEPCLDQPLIVWE 276

Db 263 GTFOKAAVVPVPSGQQRVTCVHQHGLPKPLTLRWE 299

RESULT 14

I38442

Gene HLA-A-0205 protein - human

C;Species: Homo sapiens (man)

C;Date: 02-Jul-1996 #sequence_revision 02-Jul-1996 #text_change 21-Jan-2000

C;Accession: I38442

R;Holmes, N.; Ennis, P.; Wan, A.M.; Denney, D.W.; Parham, P.

J. Immunol. 139, 936-941, 1987

A;Title: Multiple genetic mechanisms have contributed to the generation of the HLA-A2/A2

A;Reference number: I38441; MUID:87252273; PMID:3496393

A;Accession: I38442

A;Status: preliminary; translated from GB/EMBL/DDBJ

A;Molecule type: DNA

A;Residues: 1-365 <RES>

A;Cross-references: EMBL:U03862; NID:9432436; PIDN:AAA03603.1; PID:9432437

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.2%; Score 503; DB 2; Length 365;
Best Local Similarity 39.7%; Pred. No. 2.5e-34;
Matches 110; Conservative 43; Mismatches 116; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASQDGLSLFEALGVYDDQLFVFDHE--SRVPRTPWSSRISSQMW 62

Db 26 SHSMRYFTTSVRGEPGFIAVGVDQTFVRFDSDAASQRMPEAPWIEQE-GPEYW 84

QY 63 LQLSLSLKGWDHMTFTVDFTIMENHNASKB-SHTLQVILGCMEQED-NSTEGYWKYGYDG 120

Db 85 DGETRKKVKAHSQTHRVDLGLTRGYNQSEAGSHTVQRMVGCVDGSDWRFLRGVHQYAYDG 144

QY 121 QDALEFCPDTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180

Db 145 KDYIALKEDLRSWTAADMAAQTTHKWEAAHV-AEQRAYLEGTCVWLRRLRYLENGKETL 203

QY 181 DQOVPLPVKVTTH-VTSSVTLTLCRALNYPQNTMKWLKDKQPMDAKFEPEKDVLPNGD 239

Db 204 QRTDAPKTHMTHAVSDHEATLRCWALSFPABITLTWQDGDG-OTQDTLVETRPAGD 262

QY 240 GTYQGHITLAVPGEQRVTCVHEPCLDQPLIVWE 276

Db 263 GTFOKAAVVPVPSGQQRVTCVHQHGLPKPLTLRWE 299

RESULT 15

I61902

MHC class I histocompatibility antigen HLA-A alpha chain precursor - human (isolate A*0.

C;Species: Homo sapiens (man)

A;Variety: isolate A*0212

C;Date: 06-Sep-1996 #sequence_revision 06-Sep-1996 #text_change 23-Jul-1999

C;Accession: I61902

R;Belich, M.P.; Madrigal, J.A.; Hildebrand, W.H.; Zemmour, J.; Williams, R.C.; Luz, R.;

Nature 357, 326-329, 1992

A;Title: Unusual HLA-B alleles in two tribes of Brazilian Indians.

A;Reference number: I37120; MUID:92269955; PMID:1317015

A;Accession: I61902

A;Status: translated from GB/EMBL/DDBJ

A;Molecule type: mRNA

A;Residues: 1-365 <RES>

A;Cross-references: GB:M84378; NID:9187625; PIDN:AAA59604.1; PID:9187626

A;Experimental source: cell line KRC 033; isolate A*0212

C;Genetics:

A;Gene: GDB:HLA-A

A;Cross-references: GDB:119310; OMIM:142800

A;Map position: 6p21.3-6p21.3

C;Superfamily: class I histocompatibility antigen; immunoglobulin homology

C;Keywords: transmembrane protein

F;1-24/Domain: signal sequence #status predicted <SIG>

F;25-365/Product: MHC class I histocompatibility antigen HLA-A alpha chain #status pred.

F;220-285/Domain: immunoglobulin homology <IMM>

Query Match 33.2%; Score 503; DB 2; Length 365;
Best Local Similarity 39.4%; Pred. No. 2.5e-34;
Matches 109; Conservative 44; Mismatches 116; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASEQDGLSLFEALGVYDDQLFVFDHE--SRVPRTPWSSRISSQMW 62

Db 26 SHSMRYFTTSVRGEPGFIAVGVDQTFVRFDSDAASQRMPEAPWIEQE-GPEYW 84

QY 63 LQLSLSLKGWDHMTFTVDFTIMENHNASKB-SHTLQVILGCMEQED-NSTEGYWKYGYDG 120

Db 85 DGETRKKVKAHSQTHRVDLGLTRGYNQSEAGSHTVQRMVGCVDGSDWRFLRGVHQYAYDG 144

QY 121 QDALEFCPDTLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180

Db 145 KDYIALKEDLRSWTAADMAAQTTHKWEAAHV-AEQRAYLEGTCVWLRRLRYLENGKETL 203

QY 181 DQOVPLPVKVTTH-VTSSVTLTLCRALNYPQNTMKWLKDKQPMDAKFEPEKDVLPNGD 239

Db 204 QRTDAPKTHMTHAVSDHEATLRCWALSFPABITLTWQDGDG-OTQDTLVETRPAGD 262

QY 240 GTYQGHITLAVPGEQRVTCVHEPCLDQPLIVWE 276

Db 263 GTFOKAAVVPVPSGQQRVTCVHQHGLPKPLTLRWE 299

Search completed: August 5, 2003, 13:10:37

Job time : 16 secs

[7] SEQUENCE FROM N.A. (ISOFORMS 1; 7; 8; 9 AND 10).
 RP MEDLINE=20448010; PubMed=11001625;
 RA Thénie A., Orhant M., Gicquel I., Fergelot P., Le Gall J.-Y.,
 RA David V., Mosser J.;
 RT "The HFE gene undergoes alternate splicing processes.";
 RL Blood Cells Mol. Dis. 26:155-162(2000).
 RN [8]
 RP FUNCTION.
 RP MEDLINE=98132614; PubMed=9465039;
 RA Feder J.N., Penny D.M., Irrinki A., Lee V.K., Lebron J.A., Watson N.,
 RA Tsuchihashi Z., Sigal E., Bjorkman P.J., Schatzman R.C.;
 RT "The hemochromatosis gene product complexes with the transferrin
 RT receptor and lowers its affinity for ligand binding.";
 RL Proc. Natl. Acad. Sci. U.S.A. 95:1472-1477(1998).
 RN [9]
 RP X-RAY CRYSTALLOGRAPHY (2.6 ANGSTROMS).
 RP MEDLINE=98206473; PubMed=9546397;
 RA Lebron J.A., Bennett M.J., Vaughn D.E., Chirino A.J., Snow P.M.,
 RA Mintier G.A., Feder J.N., Bjorkman P.J.;
 RT "Crystal structure of the hemochromatosis protein HFE and
 RT characterization of its interaction with transferrin receptor.";
 RL Cell 93:111-123(1998).
 RN [10]
 RP VARIANTS HH ASP-63 AND TYR-282.
 RP MEDLINE=97260408; PubMed=9106528;
 RA Carella M., D'Ambrosio L., Totaro A., Grifa A., Valentino M.A.,
 RA Piperno A., Girelli D., Roetto A., Franco B., Gasparini P.,
 RA Camaschella C.;
 RT "Mutation analysis of the HLA-H gene in Italian hemochromatosis
 RT patients.";
 RL Am. J. Hum. Genet. 60:828-832(1997).
 RN [11]
 RP VARIANT HH/PCT TYR-282.
 RP MEDLINE=97176837; PubMed=9024376;
 RA Roberts A.G., Whitley S.D., Morgan R.R., Worwood M., Elder G.H.;
 RT "Increased frequency of the hemochromatosis Cys282Tyr mutation in
 RT sporadic porphyria cutanea tarda.";
 RL Lancet 349:321-323(1997).
 RN [12]
 RP VARIANTS HH/PCT ASP-63.
 RP MEDLINE=98085904; PubMed=9425935;
 RA Sampietro M., Piperno A., Lupica L., Arosio C., Vergani A.,
 RA Corbetta N., Malosio I., Mattioli M., Fracanzani A.L.,
 RA Cappellini M.D., Fiorelli G., Fargion S.;
 RT "High prevalence of the H163Asp HFE mutation in Italian patients with
 RT porphyria cutanea tarda.";
 RL Hepatology 27:181-184(1998).
 RN [13]
 RP VARIANTS HH/PCT ASP-63 AND TYR-282.
 RP MEDLINE=98281650; PubMed=9620340;
 RA Bonkovsky H.L., Poh-Fitzpatrick M., Pimstone N., Obando J.,
 RA Di Bisceglie A., Tattire C., Tortorelli K., LeClair P., Mercurio M.G.,
 RA Lambercht R.W.;
 RT "Porphyria cutanea tarda, hepatitis C, and HFE gene mutations in North
 RT America.";
 RL Hepatology 27:1661-1669(1998).
 RN [14]
 RP VARIANTS HH ASP-63, CYS-65 AND TYR-282.
 RP MEDLINE=99211934; PubMed=10194428;
 RA Mura C., Ragues O., Ferec C.;
 RT "HFE mutations analysis in 711 hemochromatosis probands: evidence for
 RT S65C implication in mild form of hemochromatosis.";
 RL Blood 93:2502-2505(1999).
 RN [15]
 RP VARIANTS HH CYS-65; ARG-93 AND THR-105.
 RP MEDLINE=20042794; PubMed=10575540;
 RA Barton J.C., Sawada-Hirai R., Rothenberg B.E., Acton R.T.;
 RT "Two novel missense mutations of the HFE gene (I105T and G93R) and
 RT identification of the S65C mutation in Alabama hemochromatosis
 RT probands.";
 RL Blood Cells Mol. Dis. 25:147-155(1999).
 RN [16]

RP VARIANTS VP ASP-63 AND HIS-127, VARIANT HH MET-330, AND VARIANTS
 RP MET-53 AND MET-59.
 RX MEDLINE=99330560; PubMed=10401000;
 RA de Villiers J.N.P., Hillermann R., Loubser L., Kotze M.J.;
 RT "Spectrum of mutations in the HFE gene implicated in haemochromatosis
 RT and porphyria.";
 RL Hum. Mol. Genet. 8:1517-1522(1999).
 RN [17]
 RP VARIANTS HH ASP-63 AND TYR-282.
 RP MEDLINE=99140360; PubMed=10094552;
 RA Merryweather-Clarke A.T., Simonsen H., Shearman J.D., Pointon J.J.,
 RA Norgaard-Pedersen B., Robson K.J.H.;
 RT "A retrospective anonymous pilot study in screening newborns for HFE
 RT mutations in Scandinavian populations.";
 RL Hum. Mutat. 13:154-159(1999).
 RN [18]
 RP VARIANT HH CYS-65.
 RP Fagan E., Payne S.J.;
 RT "A novel missense mutation S65C in the HFE gene with a possible role
 RT in hereditary haemochromatosis.";
 RL Hum. Mutat. 13:507-508(1999).
 RN [19]
 RP VARIANT LYS-277.
 RP MEDLINE=20081073; PubMed=10612845;
 RA Bradbury R., Fagan E., Payne S.J.;
 RT "Two novel polymorphisms (E277K and V212V) in the haemochromatosis
 RT gene HFE.";
 RL Hum. Mutat. 15:120-120(2000).
 CC -!- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
 CC affinity for iron-loaded transferrin.
 CC -!- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -!- ALTERNATIVE PRODUCTS:
 CC Event=Alternative splicing; Named isoforms=10;
 CC Comment=Additional isoforms seem to exist;
 CC Name=1;
 CC IsoId=Q30201-1; Sequence=Displayed;
 CC Name=2; Synonyms=delE2;
 CC IsoId=Q30201-2; Sequence=VSP_003218;
 CC Name=3; Synonyms=del14B4;
 CC IsoId=Q30201-3; Sequence=VSP_003225;
 CC Name=4; Synonyms=delE214E4;
 CC IsoId=Q30201-4; Sequence=VSP_003219, VSP_003225;
 CC Name=5;
 CC IsoId=Q30201-5; Sequence=VSP_003219;
 CC Name=6;
 CC IsoId=Q30201-6; Sequence=VSP_003220;
 CC Name=7; Synonyms=delE3;
 CC IsoId=Q30201-7; Sequence=VSP_003221;
 CC Name=8; Synonyms=1043-2283del.introns6;
 CC IsoId=Q30201-8; Sequence=VSP_003226, VSP_003227;
 CC Name=9; Synonyms=delE3-7;
 CC IsoId=Q30201-9; Sequence=VSP_003223, VSP_003224;
 CC Name=10; Synonyms=562-878del;
 CC IsoId=Q30201-10; Sequence=VSP_003222;
 CC -!- TISSUE SPECIFICITY: IN ALL TISSUES TESTED EXCEPT BRAIN.
 CC -!- DISEASE: DEFECTS IN HFE ARE A CAUSE OF HEREDITARY HEMOCHROMATOSIS
 CC (HH). HH IS AN AUTOSOMAL RECESSIVE INBORN DISORDER OF IRON
 CC METABOLISM. FREQUENT AMONG CAUCASIANS. HH IS CHARACTERIZED BY
 CC ABNORMAL INTESTINAL IRON ABSORPTION AND PROGRESSIVE INCREASE OF
 CC TOTAL BODY IRON, WHICH RESULTS IN MIDLIFE IN CLINICAL
 CC COMPLICATIONS INCLUDING CIRRHOSIS, CARDIOPATHY, DIABETES,
 CC ENDOCRINE DYSFUNCTIONS, ARTHROPATHY, AND SUSCEPTIBILITY TO LIVER
 CC CANCER. SINCE THE DISEASE COMPLICATIONS CAN BE EFFECTIVELY
 CC PREVENTED BY REGULAR PHLEBOTOMIES, EARLY DIAGNOSIS IS MOST
 CC IMPORTANT TO PROVIDE A NORMAL LIFE EXPECTANCY TO THE AFFECTED
 CC SUBJECTS.
 CC -!- DISEASE: DEFECTS IN HFE ARE A CAUSE OF PORPHYRIA CUTANEA TARDA
 CC (PCT), A DISORDER CHARACTERIZED BY LIGHT-SENSITIVE DERMATITIS AND
 CC PRESENCE OF LARGE AMOUNTS OF UROPORPHYRIN IN URINE. IRON OVERLOAD
 CC IS OFTEN PRESENT IN ASSOCIATION WITH VARYING DEGREES OF LIVER
 CC DAMAGE. PCT IS THE MOST COMMON FORM OF PORPHYRIA WORLDWIDE. IT
 CC OCCURS IN TWO FORMS: THE SPORADIC TYPE (PCT TYPE I) AND THE
 CC FAMILIAL TYPE (PCT TYPE II), WHICH IS INHERITED IN AN AUTOSOMAL

Query Match 99.2%; Score 1502; DB 1; Length 348;
Best Local Similarity 99.3%; Pred. No. 2,3e-118;
Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDDQLFVYDHSRRVPRTPWSSRISQ 60
DB 23 RLLRSHSLHYLFMGASQDGLSLFEALGYVDDQLFVYDHSRRVPRTPWSSRISQ 82
QY 61 MWLQSLQSLKGDHMFVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGDHMFVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDLEFCPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQVPPPLVKKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKVLPNGDG 240
DB 203 DQVPPPLVKKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKVLPNGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 298

RESULT 2
HFE DICSU
ID HFE DICSU STANDARD; PRT; 348 AA.
AC Q9GLA2;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Hereditary hemochromatosis protein precursor.
GN HFE.
OS Dicerorhinus sumatrensis (Sumatran rhinoceros).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Perissodactyla; Rhinocerotidae; Dicerorhinus.
OX NCBI_TaxID=89632;
[1]
SEQUENCE FROM N.A.
RA West C.J., Worley M., Beutler E.;
RT "Rhinoceros HFE polymorphisms."
RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
CC affinity for iron-loaded transferrin.
CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
CC -1- SIMILARITY: TO MHC CLASS I ANTIGENS.
CC
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AY007543; AAG23703.1; --
DR HSP; Q30201; 1A6Z.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig-cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig_1.
DR Pfam; PF00129; MHC_I; 1.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGC1; 1.
DR PROSITE; PSS0835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
DR MHC_I; Transmembrane; Glycoprotein; Signal.
KW MHC_I; 22 BY SIMILARITY.
FT SIGNAL 1 22
FT CHAIN 23 348 HEREDITARY HEMOCHROMATOSIS PROTEIN.
FT DOMAIN 23 114 EXTRACELLULAR ALPHA-1.

FT DOMAIN 115 205 EXTRACELLULAR ALPHA-2.
FT DOMAIN 206 297 EXTRACELLULAR ALPHA-3.
FT TRANSMEM 298 307 CONNECTING PEPTIDE.
FT TRANSMEM 307 330 POTENTIAL.
FT DOMAIN 331 348 CYTOPLASMIC TAIL.
FT DISULFID 124 187 BY SIMILARITY.
FT DISULFID 225 282 BY SIMILARITY.
FT CARBOHYD 110 110 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 130 130 N-LINKED (GLCNAC. .) (POTENTIAL).
FT CARBOHYD 234 234 N-LINKED (GLCNAC. .) (POTENTIAL).
SQ SEQUENCE 348 AA; 39740 MW; 518BFD357AB83B90 CRC64;

Query Match 81.0%; Score 1227; DB 1; Length 348;
Best Local Similarity 81.0%; Pred. No. 2.2e-95;
Matches 221; Conservative 20; Mismatches 32; Indels 0; Gaps 0;

QY 4 RSHSLHYLFMGASQDGLSLFEALGYVDDQLFVYDHSRRVPRTPWSSRISQMWL 63
DB 26 RSHSLHYLFMGASQDGLSLFEALGYVDDQLFVYDHSRRVPRTPWSSRISQMWL 85
QY 64 QLSQSLKGDHMFVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDGQA 123
DB 86 QLSQSLKGDHMFVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDGQ 145
QY 124 LSFCDPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 183
DB 146 LSFCDPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 205
QY 184 VPPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKVLPNGDGTQ 243
DB 206 VPPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKVLPNGDGTQ 265
QY 244 GWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
DB 266 SWVALAVPPGEEQRYTCQVEHPGLDQPLIVWE 298

RESULT 3
HFE CERSI
ID HFE CERSI STANDARD; PRT; 348 AA.
AC Q9GKZ0;
DT 28-FEB-2003 (Rel. 41, Created)
DT 28-FEB-2003 (Rel. 41, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Hereditary hemochromatosis protein precursor.
GN HFE.
OS Ceratotherium simum (White rhinoceros) (Square-lipped rhinoceros).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Perissodactyla; Rhinocerotidae; Ceratotherium.
OX NCBI_TaxID=9807;
[1]
SEQUENCE FROM N.A.
RA West C.J., Worley M., Beutler E.;
RT "Rhinoceros HFE polymorphisms."
RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
CC affinity for iron-loaded transferrin.
CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
CC -1- SIMILARITY: TO MHC CLASS I ANTIGENS.
CC
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL; AY007541; AAG23701.1; --
DR HSP; Q30201; 1A6Z.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig-cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig_1.
DR Pfam; PF00129; MHC_I; 1.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGC1; 1.
DR PROSITE; PSS0835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
DR MHC_I; Transmembrane; Glycoprotein; Signal.
KW MHC_I; 22 BY SIMILARITY.
FT SIGNAL 1 22
FT CHAIN 23 348 HEREDITARY HEMOCHROMATOSIS PROTEIN.
FT DOMAIN 23 114 EXTRACELLULAR ALPHA-1.

OX NCBI_TaxID=9805;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA West C.J., Worley M., Beutler E.;
 RT "Rhinceros HFE polymorphisms";
 RL Submitted (AUG-2000) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
 CC affinity for iron-loaded transferrin.
 CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -1- SIMILARITY: TO MHC CLASS I ANTIGENS.
 CC -----
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 CC -----
 DR EMBL; AY007542; AAG23702.1; -;
 DR EMBL; AF301592; AAG39940.1; -;
 DR HSSP; Q30201; 1A62.
 DR InterPro; IPR007110; Ig-like.
 DR InterPro; IPR003597; Ig-cl.
 DR InterPro; IPR003006; Ig_MHC.
 DR InterPro; IPR001039; MHC_I.
 DR Pfam; PF00047; Ig; 1.
 DR Pfam; PF00129; MHC_I; 1.
 DR ProDom; PD000050; MHC_I; 1.
 DR SMART; SM00407; Igc1; 1.
 DR PROSITE; PS50835; IG_LIKE; 1.
 DR PROSITE; PS00290; IG_MHC; 1.
 DR MHC I; Transmembrane; Glycoprotein; Signal.
 FT SIGNAL 1 22 BY SIMILARITY.
 FT CHAIN 23 348 HEREDITARY HEMOCHROMATOSIS PROTEIN.
 FT DOMAIN 23 114 EXTRACELLULAR ALPHA-1.
 FT DOMAIN 115 205 EXTRACELLULAR ALPHA-2.
 FT DOMAIN 206 297 EXTRACELLULAR ALPHA-3.
 FT DOMAIN 298 306 CONNECTING PEPTIDE.
 FT TRANSMEM 307 330 POTENTIAL.
 FT DOMAIN 331 348 CYTOPLASMIC TAIL.
 FT DISULFID 124 187 BY SIMILARITY.
 FT DISULFID 225 282 BY SIMILARITY.
 FT CARBOHYD 110 110 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 130 130 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 234 234 N-LINKED (GLCNAC. . .) (POTENTIAL).
 SQ SEQUENCE 348 AA; 39818 MW; 4D95E7B01E48FB90 CRC64;
 Query Match 80.4%; Score 1218; DB 1; Length 348;
 Best Local Similarity 80.6%; Pred. No. 1.2e-94;
 Matches 220; Conservative 20; Mismatches 33; Indels 0; Gaps 0;
 QY 4 RSHSLHYLFGASEQDLGLSLFALGVDDQLFVFDHESRRVPEPTPWSSRISSQWL 63
 DB 26 RSHSLRYLFGASERDGLFLFALGVDDQLFVFDHESRRVPEPTPWSSRISSQWL 85
 QY 64 QLSQSLKGDHMTFVDFWTIMENHNASKBSHTLQVILGCEMQRDSTEGWKYGYDQDA 123
 DB 86 QLTQSLKGDHMTFVDFWTIMDNHNHKSHTLQVILGCEVQBDNSTGRFWKYGQDQH 145
 QY 124 LEFCPTDLWRRAEPRAPWPTKLEWRHKIRARQNRAYLERDCPAQLQLLELGRVLDQ 183
 DB 146 LEFCPTDLWRRAESRALTTKLEWEVKNIRAKQNRAYLERDCPEQLQLLELGRVLDQ 205
 QY 184 VPLVVKTHVTSVTLRCRALNYPTNTMKWLKDKQPMADKEPEPKDVLNPGNGTGY 243
 DB 206 VPLVVKTHVTSVTLRCQALNFPQNTITRWLKDKEVDVVDKAEKDVLPVSGDGTGY 265
 QY 244 GWITLAVPPGEORYTCVESHGGLDQPLVIWE 276
 DB 266 SWEALAVPPGEORYTCVESHGGLDQPLVIWE 298

RESULT 6
 HFE RAT
 ID HFE RAT STANDARD; PRT; 360 AA.
 AC O35799; O35175;
 DT 15-JUL-1998 (Rel. 36, Created)
 DT 15-JUL-1998 (Rel. 36, Last sequence update)
 DT 28-FEB-2003 (Rel. 41, Last annotation update)
 DE Hereditary hemochromatosis protein homolog precursor (RT1-CAFE).
 GN HFE.
 OS Rattus norvegicus (Rat).
 OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Rattus.
 OC NCBI_TaxID=10116;
 RN [1]
 RP SEQUENCE FROM N.A.
 RA Banasch M.W., Schaefer H., Schmidt W.E.;
 RL Submitted (SEP-1997) to the EMBL/GenBank/DBJ databases.
 RN [2]
 RP SEQUENCE OF 42-303 FROM N.A.
 RC TISSUE=Small intestine;
 RA Sawada-Hirai R., Rothenberg B.E.;
 RL Submitted (JUN-1997) to the EMBL/GenBank/DBJ databases.
 CC -1- FUNCTION: BINDS TO TRANSFERRIN RECEPTOR (TFR) AND REDUCES ITS
 CC AFFINITY FOR IRON-LOADED TRANSFERRIN (BY SIMILARITY).
 CC -1- SUBCELLULAR LOCATION: Type I membrane protein.
 CC -1- SIMILARITY: TO MHC CLASS I ANTIGENS.
 CC -----
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 CC -----
 DR EMBL; AJ001517; CAA04799.1; -;
 DR EMBL; AF008587; AAB86597.1; -;
 DR HSSP; Q30201; 1A62.
 DR InterPro; IPR007110; Ig-like.
 DR InterPro; IPR003597; Ig-cl.
 DR InterPro; IPR003006; Ig_MHC.
 DR InterPro; IPR001039; MHC_I.
 DR Pfam; PF00047; Ig; 1.
 DR Pfam; PF00129; MHC_I; 1.
 DR PRINTS; PR01638; MHCCLASSI.
 DR ProDom; PD000050; MHC_I; 1.
 DR SMART; SM00407; Igc1; 1.
 DR PROSITE; PS50835; IG_LIKE; 1.
 DR PROSITE; PS00290; IG_MHC; 1.
 DR MHC I; Transmembrane; Glycoprotein; Signal.
 FT SIGNAL 1 25 POTENTIAL.
 FT CHAIN 26 360 HEREDITARY HEMOCHROMATOSIS PROTEIN
 FT DOMAIN 26 127 HOMOLOG.
 FT DOMAIN 128 218 EXTRACELLULAR ALPHA-1.
 FT DOMAIN 219 310 EXTRACELLULAR ALPHA-2.
 FT DOMAIN 311 319 EXTRACELLULAR ALPHA-3.
 FT TRANSMEM 320 340 CONNECTING PEPTIDE.
 FT DOMAIN 341 360 POTENTIAL.
 FT DISULFID 137 200 CYTOPLASMIC TAIL.
 FT DISULFID 238 295 BY SIMILARITY.
 FT CARBOHYD 115 115 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 143 143 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 167 167 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CARBOHYD 247 247 N-LINKED (GLCNAC. . .) (POTENTIAL).
 FT CONFLICT 198 198 R -> K (IN REF. 2).
 SQ SEQUENCE 360 AA; 40988 MW; CC819834EB3240B3 CRC64;
 Query Match 75.6%; Score 1145; DB 1; Length 360;
 Best Local Similarity 73.2%; Pred. No. 1.6e-88;
 Matches 205; Conservative 29; Mismatches 38; Indels 8; Gaps 1;

```
Qy 5 SHSLHYLFMGASEODLGLSFEALGYVDDQLFVYDHSRRVPRTPWVSSRISSQWLQ 64
Db 32 SHSLRYLFMGASRPDLGLPFEALGYVDDQLFVSYNHSRAEPRAPIWILQOTSSQLWLQ 91
Qy 65 LSQSLKGWDMFTVDFTIMENHNASK-----ESHTLQVILGCEMQEDNSTEGYWKY 116
Db 92 LSQSLKGWDMFTVDFTIMENHNASK-----ESHTLQVILGCEMQEDNSTEGYWKY 151
Qy 117 GYGDQDALEFCPDTLDWRAAEPRAPWPKLEWERHKIRARONRAYLERDCPAQLQQLLELG 176
Db 152 GYGDQDHLEFCPKTLNWSAAEPRAWATMEWEHRIRAROSRYLQDCPQQQQLVLELQ 211
Qy 177 RGVLDDQVPLVKVTHVTSSVTLRCALNYPQNTIMKWLKDQKPMDAKEFEKDVLP 236
Db 212 RGVLGQVPTLVKVTREHASTGTSLRCQALNFFPQNTIMRWLXDSQPLDAKDVNPENLVP 271
Qy 237 NGDGTGQWTLAVPGEORVYTCQVEHPGLDQPLVIWE 276
Db 272 NGDGTGQWTLAVAPGEETRFSCQVEHPGLDQPLATWE 311

RESULT 7
HFE_MOUSE STANDARD; PRT; 359 AA.
AC P70387;
DT 15-JUL-1998 (Rel. 36, Created)
DT 15-JUL-1998 (Rel. 36, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE Hereditary hemochromatosis protein homolog precursor.
GN HFE OR MR2.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]
RN SEQUENCE FROM N.A.
RC STRAIN=129/SvJ;
RC MDLINE=98060831; PubMed=9396865;
RA Riegert P., Gilfillan S., Nanda I., Schmid M., Bahram S.;
RT "the mouse HFE gene.";
RL Immunogenetics 47:174-177 (1998).
RN [2]
RN SEQUENCE FROM N.A.
RC STRAIN=BALB/c; TISSUE=Lung;
RA Hashimoto K.;
RL Submitted (SEP-1996) to the EMBL/GenBank/DBJ databases.
RN [3]
RN SEQUENCE OF 37-211 FROM N.A.
RC STRAIN=BALB/c; TISSUE=Liver;
RC MDLINE=97148566; PubMed=9020055;
RA Hashimoto K., Hirai M., Kurosawa Y.;
RT "Identification of a mouse homolog for the human hereditary
hemochromatosis candidate gene.";
RL Biochem. Biophys. Res. Commun. 230:35-39 (1997).
RN [4]
RN SEQUENCE OF 79-359 FROM N.A.
RC STRAIN=129;
RA Albright W., Drabant B., Doenecke D.;
RL Submitted (MAY-1997) to the EMBL/GenBank/DBJ databases.
CC -I- FUNCTION: Binds to transferrin receptor (TFR) and reduces its
affinity for iron-loaded transferrin (By similarity).
CC -I- SUBCELLULAR LOCATION: Type I membrane protein.
CC -I- SIMILARITY: TO MHC CLASS I ANTIGENS.
CC
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CC -----
CC EMBL; AF007558; AAC03447.1; --
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DR EMBL; U66849; AAB07525.1; --
DR EMBL; Y12650; CAA73197.1; --
DR EMBL; U80604; AAB51504.1; --
DR PIR; JC5382; JC5382.
DR HSSP; Q30201; IA62.
DR MGD; MGI:109191; Hfe.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig_c1.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGC1; 1.
DR PROSITE; PS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal.
FT SIGNAL 1 24
FT CHAIN 25 359
FT HEREDITARY HEMOCHROMATOSIS PROTEIN
FT HOMOLOG.
FT DOMAIN 25 126
FT DOMAIN 127 217
FT DOMAIN 218 309
FT DOMAIN 310 318
FT TRANSMEM 319 339
FT DOMAIN 340 359
FT DISULFID 136 139
FT DISULFID 237 294
FT CARBOHYD 114 114
FT CARBOHYD 142 142
FT CARBOHYD 166 166
FT CARBOHYD 246 246
FT SEQUENCE 359 AA; 40548 MW; 4BDE6C27F9FF20B4 CRC64;
SQ
Query Match 74.6%; Score 1129; DB 1; Length 359;
Best Local Similarity 71.9%; Pred. No. 3.5e-87;
Matches 202; Conservative 30; Mismatches 41; Indels 8; Gaps 1;
Qy 4 RSHSLHYLFMGASEODLGLSFEALGYVDDQLFVYDHSRRVPRTPWVSSRISSQWL 63
Db 30 RSHSLRYLFMGASEPDLGLPFEALGYVDDQLFVSYNHSRAEPRAPIWILQOTSSQL 89
Qy 64 QLSQSLKGWDMFTVDFTIMENHNASK-----ESHTLQVILGCEMQEDNSTEGYWK 115
Db 90 HLSQSLKGWDMFTVDFTIMENHNASK-----ESHTLQVILGCEMQEDNSTEGYWK 149
Qy 116 YGYDQDALEFCPDTLDWRAAEPRAPWPKLEWERHKIRARONRAYLERDCPAQLQQLLEL 175
Db 150 YGYDQDHLEFCPKTLNWSAAEPRAWATMEWEHRIRAROSRYLQDCPQQQQLVLEL 209
Qy 176 GYGVLDQVPLVKVTHVTSSVTLRCALNYPQNTIMKWLKDQKPMDAKEFEKDVLP 235
Db 210 GYGVLDQVPTLVKVTREHASTGTSLRCQALNFFPQNTIMRWLXDSQPLDAKDVNPEN 269
Qy 236 PNGDGTGQWTLAVPGEORVYTCQVEHPGLDQPLVIWE 276
Db 270 PNGDGTGQWTLAVAPGEETRFSCQVEHPGLDQPLATWE 310

RESULT 8
HAIA_RABIT STANDARD; PRT; 361 AA.
ID HAIA_RABIT
AC P01894;
DT 21-JUL-1986 (Rel. 01, Created)
DT 21-JUL-1986 (Rel. 01, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE RIA class I histocompatibility antigen, alpha chain 11/11 precursor.
OS Oryctolagus cuniculus (Rabbit).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Lagomorpha; Leporidae; Oryctolagus.
OX NCBI_TaxID=9986;
RN [1]
```


QY 121 QDALEFCPDTLDWRAAEPRAPWTKLEWHRKIRARONRAYLERDCPAQLQQLLELGRGVL 180
DB 145 ADYIALNEDLRSWTAAQTQKWEAAG-EAERHAYLERECVWELRYLEMGKCTL 203
QY 181 DQVPPPLVKVTHVTS-VTTLRCRALNYPQNTMKWLKDKQPMDAKEFEKPKDVLNPGD 239
DB 204 QRADPPKRAVTHHPASDREATLRCWALGYPABEITLTWQDGED-QTQDTVELVETRPAGD 262
QY 240 GTYQGWITLAVPGEORVTCOVHEPGLDPLIVINE 276
DB 263 GTFOKAAVVPVSGEORVTCOVHEPGLDPLIVINE 299

RESULT 10

ID _IA01 PANTR STANDARD; PRT; 365 AA.
AC P16209;
DT 01-APR-1990 (Rel. 14, Created)
DT 01-APR-1990 (Rel. 14, Last sequence update)
DT 01-APR-1993 (Rel. 25, Last annotation update)
DE CHLA class I histocompatibility antigen, A-2 alpha chain precursor.
DE Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
OX NCBI_TaxID=9598;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=90201944; PubMed=1690682;
RA Lawlor D.A., Warren E., Ward F.E., Parham P.;
RT "Comparison of class I MHC alleles in humans and apes.";
RL Immunol. Rev. 113:147-185(1990).
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
CC THE IMMUNE SYSTEM.
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN).
CC
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL; M30678; AAA7970.1; -;
CC PIR; I36961; I36961.
CC HSP; Q95352; IHKK.
CC InterPro; IPR007110; Ig-like.
CC InterPro; IPR003597; Ig cl.
CC InterPro; IPR003006; Ig_MHC.
CC InterPro; IPR001039; MHC_I.
CC Pfam; PF00047; Ig; 1.
CC Pfam; PF00129; MHC_I; 1.
CC PRINTS; PR01638; MHCCLASS1.
CC ProDom; PD000050; MHC_I; 1.
CC SMART; SM00407; IGc1; 1.
CC PROSITE; PS50835; IG_LIKE; 1.
CC PROSITE; PS00290; IG_MHC; 1.
CC MHC I; Transmembrane; Glycoprotein; Signal.
CC SIGNAL 1 24
CC CHAIN 25 365

CHLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
A-2 ALPHA CHAIN.
DOMAIN 25 114
EXTRACELLULAR ALPHA-1.
DOMAIN 115 206
EXTRACELLULAR ALPHA-2.
DOMAIN 207 298
EXTRACELLULAR ALPHA-3.
DOMAIN 299 308
CONNECTING PEPTIDE.
DOMAIN 309 332
DOMAIN 333 365
CYTOPLASMIC TAIL.
DOMAIN 125 188
BY SIMILARITY.
DISULFID 227 283
BY SIMILARITY.
FT CARBOHYD 110 110
N-LINKED (GLCNAC...) (BY SIMILARITY).
SQ SEQUENCE 365 AA; 40848 MW; FC452786BD038D3E CRC64;

Query Match 33.8%; Score 511; DB 1; Length 365;
Best Local Similarity 39.7%; Pred. No. 1.6e-35;
Matches 110; Conservative 44; Mismatches 115; Indels 8; Gaps 7;
QY 5 SHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHE--SRVPERPTWVSSRISQW 62
DB 26 SHSMRYFFTSVSPGCGEPRTIAVGVDVDTQVRFDSDAASQRMPEPRAPWIEQE-GPEYW 84
QY 63 LQLSLSLKGDWHMFTVDFWTIMENNASKE-SHTLQVILGCEMQEDNS--TEGYWYGYDG 120
DB 85 DEETRSKSAHSQTDVRDVLGTLRGYNQSDGSHITQIMYGCDVSDGRFLRGTRDAYDG 144
QY 121 QDALEFCPDTLDWRAAEPRAPWTKLEWHRKIRARONRAYLERDCPAQLQQLLELGRGVL 180
DB 145 ADYIALNEDLRSWTAAQTQKWEAAG-EAERHAYLERECVWELRYLEMGKCTL 203
QY 181 DQVPPPLVKVTHH-VTSSVTTLRCRALNYPQNTMKWLKDKQPMDAKEFEKPKDVLNPGD 239
DB 204 QRTDPPKTHHTHPISTHEATLRCWALGYPABEITLTWQDGED-QTQDTVELVETRPAGD 262
QY 240 GTYQGWITLAVPGEORVTCOVHEPGLDPLIVINE 276
DB 263 GTFOKAAVVPVSGEORVTCOVHEPGLDPLIVINE 299
RESULT 11
HAIB BOVIN STANDARD; PRT; 364 AA.
ID _HAIB BOVIN STANDARD; PRT; 364 AA.
AC P13753;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE BOLA class I histocompatibility antigen, alpha chain BL3-7 precursor.
OS Bos taurus (Bovine).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Cetartiodactyla; Ruminantia; Pecora; Bovidae;
OC Bovidae; Bovinae; Bos.
OX NCBI_TaxID=9913;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=88258075; PubMed=3133413;
RA Ennis P.D., Jackson A.P., Parham P.;
RT "Molecular cloning of bovine class I MHC cDNA.";
RL J. Immunol. 141:642-651(1988).
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
CC THE IMMUNE SYSTEM.
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN).
CC
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CC or send an email to license@isb-sib.ch).
CC
CC EMBL; M21043; AAA30641.1; -;
CC HSP; P16391; IED3.
CC InterPro; IPR007110; Ig-like.
CC InterPro; IPR003597; Ig cl.
CC InterPro; IPR003006; Ig_MHC.
CC InterPro; IPR001039; MHC_I.
CC Pfam; PF00047; Ig; 1.
CC Pfam; PF00129; MHC_I; 1.
CC PRINTS; PR01638; MHCCLASS1.
CC ProDom; PD000050; MHC_I; 1.
CC SMART; SM00407; IGc1; 1.
CC PROSITE; PS50835; IG_LIKE; 1.
CC PROSITE; PS00290; IG_MHC; 1.
CC MHC I; Transmembrane; Glycoprotein; Signal.
CC SIGNAL 1 27
CC CHAIN 28 364
BOLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT


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FT DOMAIN 28 117 ALPHA CHAIN BL3-7.
FT DOMAIN 118 209 EXTRACELLULAR ALPHA-1.
FT DOMAIN 210 301 EXTRACELLULAR ALPHA-2.
FT DOMAIN 302 310 EXTRACELLULAR ALPHA-3.
FT TRANSMEM 311 331 CONNECTING PEPTIDE.
FT DOMAIN 332 364 CYTOPLASMIC.
FT CARBOHYD 106 106 N-LINKED (GLCNAC. . .) (POTENTIAL).
FT CARBOHYD 113 113 N-LINKED (GLCNAC. . .) (BY SIMILARITY).
FT DISULFID 128 191 BY SIMILARITY.
FT DISULFID 230 286 BY SIMILARITY.
SQ SEQUENCE 364 AA; 41513 MW; 622056CF7DCFF7873 CRC64;

Query Match 33.7%; Score 510; DB 1; Length 364;
Best Local Similarity 38.9%; Pred. No. 1.9e-35;
Matches 109; Conservative 49; Mismatches 114; Indels 8; Gaps 7;

QY 2 LRSLSHLVFMGASEODLGLSLFEALGYVDDQLFVFDYDHE--SRRVERPTWVSSRISS 59
DB 26 LAGSHSLRYPTGYVSRPGLGEPRFIAVGYVDDTQFVRFDSADPNPREPRVPMWEOE-GP 84
QY 60 QMWLQLSLSKGDHMFVDFWTIMENHNASKE-SHTLQVILGCEMOEDNS-TEGYWKYG 117
DB 85 EYDWRNTRIYKDTAIFRVDLNTLRGYNQSGTSHNIQAMYGCDVGPDCGLLRGFQFG 144
QY 118 YDQDLEFCPTDLWRRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGR 177
DB 145 YDGRDVIALLNEELRSWTAADTAQAITKRWKAAAG-AAETWRNLYEGECVWELRRYLENGK 203
QY 178 GVLDQVQVPLVKVTHH-VTSVTVLRCALNYFQNTIMKLDKQPMDAKEPEPKDVL 236
DB 204 DTLRADPPKAVHTHSISDREVTLCRWALGFYPEISLWQREGD-QTDMLVELVETRP 262
QY 237 NGDGTQGWITLAVPGEQRYTCQVEHGLDQPLIVIE 276
DB 263 SGGDTQKWAALVPSGEQRYTCRVOHGLQSLPLRLWE 302

RESULT 12
ID 1A11 HUMAN STANDARD; PRT; 365 AA.
AC P13746;
DT 01-JAN-1990 (Rel. 13, Created)
DT 01-JAN-1990 (Rel. 13, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE HLA class I histocompatibility antigen, A-11 alpha chain precursor.
GN HLA-A OR HLA-A.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A. (A*1101/A*1102).
RX MEDLINE=89030641; PubMed=2460344;
RA Mayer W.E., Jonker M., Klein D., Ivanyi P., van Seventer G.,
RA Klein J.
RT "Nucleotide sequences of chimpanzee MHC class I alleles: evidence for
RT trans-species mode of evolution.";
RL EMBO J. 7:2765-2774(1988).
RN [2]
RP SEQUENCE FROM N.A. (A*1101/A*1102).
RX MEDLINE=94287401; PubMed=8016845;
RA Lin L., Tokunaga K., Ishikawa Y., Bannai M., Kashiwase K.,
RA Kuwata S., Akaza T., Tadokoro K., Shibata Y., Juji T.
RT "Sequence analysis of serological HLA-A11 split antigens, A11.1 and
RT A11.2.";
RL Tissue Antigens 43:78-82(1994).
RN [3]
RP SEQUENCE OF 26-365 FROM N.A. (A*1101).
RX MEDLINE=87192928; PubMed=2437024;
RA Cowan E.P., Jelachich M.L., Biddison W.E., Coligan J.E.;
RT "DNA sequence of HLA-A11: remarkable homology with HLA-A3 allows
RT identification of residues involved in epitopes recognized by
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RT antibodies and T cells.";
RL Immunogenetics 25:241-250(1987).
CC -I- FUNCTION: Involved in the presentation of foreign antigens to
CC the immune system.
CC -I- SUBUNIT: Dimer of alpha chain and a beta chain (beta-2-
CC microglobulin).
CC -I- POLYMORPHISM: THE FOLLOWING ALLELES OF A-11 ARE KNOWN: A*1101 (A-
CC 11E) AND A*1102 (A-11K). THE SEQUENCE SHOWN IS THAT OF A*1101.
CC -----
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CC -----
CC EMBL; X13111; CAA31503.1; --
CC EMBL; X13112; CAA31504.1; --
CC EMBL; D16841; BAA04117.1; --
CC EMBL; D16842; BAA04118.1; --
CC EMBL; M16010; AAA65449.1; --
CC EMBL; M16007; AAA65449.1; JOINED.
CC EMBL; M16008; AAA65449.1; JOINED.
CC EMBL; M16009; AAA65449.1; JOINED.
CC PIR; I83063; I83063.
CC PIR; S03536; A47636.
CC HSSP; O19673; 1HSB.
CC MIM; 142800; --
CC GO; GO:0005887; C:integral to plasma membrane; NAS.
CC GO; GO:0030106; F:MHC class I receptor activity; NAS.
CC GO; GO:0006955; P:immune response; NAS.
CC InterPro; IPR007110; IG-like.
CC InterPro; IPR003597; IG_c1.
CC InterPro; IPR003006; IG_MHC.
CC InterPro; IPR001039; MHC_I.
CC Pfam; PF00047; IG_1.
CC Pfam; PF00129; MHC_I_1.
CC PRINTS; PR01638; MHCCLASSI.
CC ProDom; PD000050; MHC_I_1.
CC SMART; SM00407; IGc1_1.
CC PROSITE; PS50835; IG_LIKE; 1.
CC PROSITE; PS00290; IG_MHC; 1.
CC MHC I; Transmembrane; Glycoprotein; Signal; Polymorphism.
FT SIGNAL 1 24
FT CHAIN 25 365 HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT DOMAIN 25 114 A-11 ALPHA CHAIN.
FT DOMAIN 115 206 EXTRACELLULAR ALPHA-1.
FT DOMAIN 207 298 EXTRACELLULAR ALPHA-2.
FT DOMAIN 299 308 EXTRACELLULAR ALPHA-3.
FT TRANSMEM 309 332 CONNECTING PEPTIDE.
FT DOMAIN 333 365 CYTOPLASMIC TAIL.
FT CARBOHYD 110 110 N-LINKED (GLCNAC. . .) (BY SIMILARITY).
FT DISULFID 125 188 BY SIMILARITY.
FT DISULFID 227 283 BY SIMILARITY.
FT VARIANT 43 43 E -> K (IN ALLELE A*1102).
FT SEQUENCE 365 AA; 40937 MW; FE49CE2D4BF6CC5 CRC64;

Query Match 33.6%; Score 508; DB 1; Length 365;
Best Local Similarity 39.4%; Pred. No. 2.8e-35;
Matches 109; Conservative 46; Mismatches 114; Indels 8; Gaps 7;

QY 5 SLSLHVLFMGASEODLGLSLFEALGYVDDQLFVFDYDHE--SRRVERPTWVSSRISSQW 62
DB 26 SLSMRVFTSVSRPGRGEPRFIAVGYVDDTQFVRFDSADPNPREPRVPMWEOE-GPEYW 84
QY 63 LOLSLSKGDHMFVDFWTIMENHNASKE-SHTLQVILGCEMOEDNS-TEGYWKYG 120
DB 85 DQETRNKQAQSDTRDVLDTLGRYNGQSGSHITQIMYGCDVGPDCGLRGRDQADVG 144
QY 121 QDALEFCPTDLWRRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRV 180
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145 KDVIALLNEDLSRTAADMMAAQTCKWEAAH-AAEQRAYLEGRCVWELRRYLENGKETL 203
181 DQOVPPVLKVTTH-VTSVVTTLRCALNYPQNTMKWLKDKQPMADAKEPEPKDVLNPGD 239
204 QRTDPKTHMTHTPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTLVELTRPAGD 262
240 GTYQGWITLAVPGGEORYTCQVEHPGLDQPLIVWE 276
263 GTFQKMAAVVPSGGEORYTCHVQHEGLPKPLTLRWE 299

RESULT 13
ID 1A03 HUMAN STANDARD; PRT; 370 AA.
AC P04439;
DT 13-AUG-1987 (Rel. 05, Created)
DT 13-AUG-1987 (Rel. 05, Last sequence update)
DT 28-FEB-2003 (Rel. 41, Last annotation update)
DE HLA class I histocompatibility antigen, A-3 alpha chain precursor.
GN HLA-A OR HLA-A.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN 1;
RP SEQUENCE FROM N.A. (A*0301).
RX MEDLINE=84207948; PubMed=6609814;
RA Strachan T., Sodoyer R., Damotte M., Jordan B.R.;
RT "Complete nucleotide sequence of a functional class I HLA gene,
RT HLA-A3: implications for the evolution of HLA genes.";
RL EMBO J. 3:887-894 (1984).
RN 2;
RP SEQUENCE FROM N.A. (A*0302).
RX MEDLINE=85290871; PubMed=2993417;
RA Cowan E.P., Jordan B.E., Colligan J.E.;
RT "Molecular cloning and DNA sequence analysis of genes encoding
RT cytotoxic T lymphocyte-defined HLA-A3 subtypes: the E1 subtype.";
RL J. Immunol. 135:2835-2841 (1985).
CC -1- FUNCTION: Involved in the presentation of foreign antigens to
CC the immune system.
CC -1- SUBUNIT: Dimer of alpha chain and a beta chain (beta-2-
CC microglobulin).
CC -1- POLYMORPHISM: THE FOLLOWING ALLELES OF A-3 ARE KNOWN: A*0301 (A-
CC 3.1) AND A*0302. THE SEQUENCE SHOWN IS THAT OF A*0301.
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CC -----
CC EMBL; X00492; CAA25162.1; ALT TERM.
DR PIR; A02192; HLHUA3.
DR HSSP; O19673; IHSB.
DR HTM; 142800;
DR GO; GO:0005887; C:integral to plasma membrane; NAS.
DR GO; GO:0030106; F:MHC class I receptor activity; NAS.
DR GO; GO:0006955; P:immune response; NAS.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_1.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal; Polymorphism.

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FT SIGNAL 1 29 HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT CHAIN 30 370 A-3 ALPHA CHAIN.
FT FT EXTRACELLULAR ALPHA-1.
FT DOMAIN 30 119 EXTRACELLULAR ALPHA-2.
FT DOMAIN 120 211 EXTRACELLULAR ALPHA-3.
FT DOMAIN 212 303 CONNECTING PEPTIDE.
FT DOMAIN 304 313
FT TRANSMEM 314 337
FT DOMAIN 338 370 CYTOPLASMIC TAIL.
FT CARBOHYD 115 115 N-LINKED (GLCNAC. .) (BY SIMILARITY).
FT DISULFID 130 193 BY SIMILARITY.
FT DISULFID 232 288 E -> V (IN ALLELE A*0302).
FT VARIANT 181 181 /FTID=VAR 004351.
FT VARIANT 185 185 L -> Q (IN ALLELE A*0302).
FT /FTID=VAR 004352.
SQ SEQUENCE 370 AA; 41368 MW; ABBIFA77460318A2 CRC64;
Query Match 33.4%; Score 506; DB 1; Length 370;
Best Local Similarity 39.6%; Pred. No. 4.2e-35;
Matches 110; Conservative 46; Mismatches 112; Indels 10; Gaps 8;

QY 5 SHSLHYLFMGASEQDGLSLFALGVDDQLFVYDHE--SRVPRTPWSSRSISQMW 62
DB 31 SHSMRYFTTSVSRPGRCGEPRFIAVGYYDDTQFVRFDSDAASQRMPEAPWIEQE-GPEYW 89
QY 63 LQLSQSLKGDHMTFTVDFTIMENNASKE-SHTLQVLGCEMQEDNS-TEGYMKYGYDG 120
DB 90 DQETRNVAQOSOTDRVDLGLTGLGYNQSEAGSHITQIMYGCVDGSDGRFLGRTRQDAYDG 149
QY 121 QDALSFPCPTLDWRAAEPRAPWPTKLEWE--RHKIRARQNRAYLERDCPAQLQLELGRGV 179
DB 150 KYIALNEDLRSWTAADMAAQITKRWAAH--AEQLRAYLDGTCEVLRRLYLENGKET 207
QY 180 LQOQVPPVLKVTTH-VTSVVTTLRCALNYPQNTMKWLKDKQPMADAKEPEPKDVLNPG 238
DB 208 LQRTDPPKTHMTHTPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTLVELTRPAG 266
QY 239 DQTYQGWITLAVPGGEORYTCQVEHPGLDQPLIVWE 276
DB 267 DGTFOKMAAVVPSGGEORYTCHVQHEGLPKPLTLRWE 304

RESULT 14
1A80 HUMAN STANDARD; PRT; 365 AA.
ID 1A80 HUMAN
AC Q09160;
DT 01-NOV-1995 (Rel. 32, Created)
DT 01-NOV-1995 (Rel. 32, Last sequence update)
DT 16-OCT-2001 (Rel. 40, Last annotation update)
DE HLA class I histocompatibility antigen, AW-80(A-1) alpha chain
DE precursor.
GN HLA-A OR HLA-A.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN 1;
RP SEQUENCE FROM N.A.
RX MEDLINE=94245293; PubMed=8198325;
RA Balas A., Garcia-Sanchez F., Gomez-Reino F., Vicario J.L.;
RT "Characterization of a new and highly distinguishable HLA-A allele in
RT a Spanish family";
RL Immunogenetics 39:452-452 (1994).
RN 2;
RP SEQUENCE FROM N.A.
RA Domena J.D.;
RL Submitted (JUN-1993) to the EMBL/GenBank/DBJ databases.
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO
CC THE IMMUNE SYSTEM.
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN).
CC -1- POLYMORPHISM: THE ONLY ALLELE OF AW-80 KNOWN IS A*8001 WHICH IS

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DR MIM; 142800; --
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig-cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW MHC I; Transmembrane; Glycoprotein; Signal; Polymorphism.
FT SIGNAL 1 24
FT CHAIN 25 365
FT HLA CLASS I HISTOCOMPATIBILITY ANTIGEN,
FT A-31 ALPHA CHAIN.
FT DOMAIN 25 114
FT DOMAIN 115 206
FT DOMAIN 207 298
FT DOMAIN 299 308
FT DOMAIN 309 332
FT TRANSMEM 333 365
FT DOMAIN 110 110
FT CARBOHYD 125 188
FT DISULFID 227 283
FT VARIANT 121 121
FT VARIANT 138 138
FT FT
FT SQ SEQUENCE 365 AA; 41004 MW; 4E760C821A3C553B CRC64;

Query Match 33.2%; Score 502; DB 1; Length 365;
Best Local Similarity 39.0%; Pred. No. 8.9e-35;
Matches 108; Conservative 50; Mismatches 111; Indels 8; Gaps 7;

QY 5 SHSLHYLFMGASQDLGLSLFEALGYDDQLFVFDHE--SRVEPRTPWVSSRISSQMW 62
   |||:| | | | | | | | | | | | | | | | | | | | | | | | | | | |
Db 26 SHSMRYFTTSVRGGEPRFIAGVYDDTQFVRFDSDAASQRMPEAPWIEQE-RPEYW 84
   | : : : | : | : | | | | | | | | | | | | | | | | | | | |
QY 63 LQLSLSLKGHDHFTVDFTWIMENHNASKE-SHTLQVILGCENQEDNS-TEGWKYGYDG 120
   | : : : | : | : | | | | | | | | | | | | | | | | | | | |
Db 85 DQETRNVAHSQIDRVDLGLTGLRGYNGSEAGSHTIQMMYCGDVGSDGRFLRGYQDDAYDG 144
   | : : : | : | : | | | | | | | | | | | | | | | | | | | |
QY 121 QDALEFCPDLDNRAEPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
   : | : | | | | | | | | | | | | | | | | | | | | | | | | |
Db 145 KDYIALNEDLRSTADMAAQITQRKEARV-AEQLRAYLEGTCTVEWLRYLENGKETL 203
   : | : | | | | | | | | | | | | | | | | | | | | | | | | |
QY 181 DQVPPPLVKVTHH-VTSSVTTLRCRALNYPQNTMKWLKDKQPMDAKEFEPKDVLPNGD 239
   : | : | | | | | | | | | | | | | | | | | | | | | | | | |
Db 204 QRTDPPKTHMTHAVSDHEATLRCWALSFPYPAITLTWRQDGED-QTQDTLVELVETRPAGD 262
   | : | : | | | | | | | | | | | | | | | | | | | | | | | |
QY 240 GTYQGWITLAVPGEORYTCQVHPGLDQPLIVINE 276
   ||:| | : | | | | | | | | | | | | | | | | | | | |
Db 263 GTFQKASVVVPSGQEQRYTCHVQHEGLPKPLTRWE 299
```

Search completed: August 5, 2003, 13:08:49
Job time : 10.5 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:05:29 ; Search time 32 Seconds
(without alignments)

2225.704 Million cell updates/sec

Title: US-10-092-404-3

Perfect score: 1514

Sequence: 1 RLRSLSLHLYFMGASEQDL.....RYTCQVEHPGLDQPLIVWE 276

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 830525 seqs, 258052604 residues

Total number of hits satisfying chosen parameters: 830525

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : SPTREMBL.23.*

- 1: sp_archaea.*
- 2: sp_bacteria.*
- 3: sp_fungi.*
- 4: sp_human.*
- 5: sp_invertebrate.*
- 6: sp_mammal.*
- 7: sp_mhc.*
- 8: sp_organelle.*
- 9: sp_phase.*
- 10: sp_plant.*
- 11: sp_rodent.*
- 12: sp_virus.*
- 13: sp_vertebrate.*
- 14: sp_unclassified.*
- 15: sp_rvirus.*
- 16: sp_bacteriap.*
- 17: sp_archaeap.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1129	74.6	358	11 Q8C2A6	Q8c2a6 mus musculus
2	1129	74.6	359	11 Q9D754	Q9d754 mus musculus
3	792	52.3	272	11 Q9R105	Q9r105 rattus norv
4	574	37.9	116	4 Q9HC69	Q9hc69 homo sapien
5	540.5	35.7	359	7 Q8HX81	Q8hx81 ornithorhyn
6	537.5	35.5	354	7 Q95HB3	Q95hb3 anas platyr
7	531.5	35.1	340	7 Q9BD50	Q9bd50 pongo pygma
8	530.5	35.0	340	7 Q9TQK3	Q9tqk3 homo sapien
9	530.5	35.0	341	4 Q9NPL2	Q9npl2 homo sapien
10	530.5	35.0	341	7 Q9B460	Q9b460 homo sapien
11	530.5	35.0	341	7 Q9BCU3	Q9bcu3 pan troglod
12	527.5	34.8	341	7 Q9BCU4	Q9bcu4 pan troglod
13	515	34.0	356	7 Q8HX66	Q8hx66 sus scrofa
14	514	33.9	332	7 Q3O990	Q3o990 pan troglod
15	514	33.9	365	7 Q9TPL7	Q9tpl7 pan troglod
16	512	33.8	105	4 Q9HC71	Q9hc71 homo sapien

ALIGNMENTS

RESULT 1

Q8C2A6 PRELIMINARY; PRT; 358 AA.

AC Q8C2A6;
DT 01-WAR-2003 (Tremblrel. 23, Created)
DT 01-WAR-2003 (Tremblrel. 23, Last sequence update)
DT 01-WAR-2003 (Tremblrel. 23, Last annotation update)
DE Hemochromatosis.
OS Mus musculus (Mouse).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
OX NCBI_TaxID=10090;
RN [1]_TaxID=10090;
RP SEQUENCE FROM N.A.
RC STRAIN=NOD; TISSUE=Thymus;
RX MEDLINE=22354683; PubMed=12466851;
RA The FANTOM Consortium,
RA the RIKEN Genome Exploration Research Group Phase I & II Team;
RT "Analysis of the mouse transcriptome based on functional annotation of
RT 60,770 full-length cDNAs."
RL Nature 420:563-573 (2002).
DR EMBL; AK088986; BAC40688.1; -.
SQ SEQUENCE 358 AA; 40421 MW; EE88FB6E5AAC844D CRC64;

Query Match 74.6%; Score 1129; DB 11; Length 358;
Best Local Similarity 71.9%; Pred. No. 1.3e-97;
Matches 202; Conservative 30; Mismatches 41; Indels 8; Gaps 1;

QY 4 RSHSLHLYFMGASEQDLGLSLFEALGYDDQLFVFDHESRVERVEPTPMVSSRISSQWL 63
Db 29 RSHSLHLYFMGASEQDLGLSLFEALGYDDQLFVFDHESRVERVEPTPMVSSRISSQWL 88
QY 64 QLSQSLKGDHNFVDFWTFIMENHNASK-----ESHTLVILGCENQDNSTEGYWK 115
Db 89 HLSQSLKGDWYMFVDFWTIMGYNHNSKVTKLGVVSESHILQVLGCEVHEDNSTSGFWR 148
QY 116 YGYDQDALLEFCPDTLDWRAAPRAWPYKLEWERHKIRARQNYLREDCPQLOLLEL 175
Db 149 YGYDQDHLFCPDKTLNWSAEPGAWATKVEWDEHKIRAKQNRDYLEKDCPQLKRLLEL 208

Qy	176	GRGVLDQQVPPLVKVTHTHVTSSVTLRCLRNLYYPQNITMKWLKDQKPMDAKEFEPKDVL	235
Db	209	GRGLVQQVPTLVKVTRHWASTGTSLRCQALDDFFQNTIMRWLNDNPDLDAKDVPNEKVL	268
Qy	236	PNGDGYTQCWITLAVPPEGEQRYTTCQVEHPGGLDPLTVIWE	276
Db	269	PNGDETYTQCVLTAVAPGDYETFTTCQVEHPGGLDPLTASWE	309


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QY 126 FCPDITLDRAAEPRAPWTKLEWRHKIRARQNRAYLERDCAQIQQLLELGRGVLDQOVP 185
DB 144 LDKDITWTTAADAQAQITKKKWEEDGTVAERRKYTYLLENTCIEWLKRKYRGKDVLEERR 203
QY 186 PLVKVTHHTVSSVTTLCRALNYYPQNTMKWLKDKQPMDAKEFEPRKDVLPNGDGTGYQGW 245
DB 204 PEVRVSGMEADKILSLSCRAHGYPFPIISLWLDGM-VQEQTQSGTVPNSDGIYHIW 262
QY 246 ITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 276
DB 263 ATIDVVPGRDKYQCRVHASLFPQGLFSWE 293

RESULT 7
Q9BD50 PRELIMINARY; PRT; 340 AA.
AC Q9BD50;
DT 01-JUN-2001 (T-EMBLrel. 17, Created)
DT 01-JUN-2001 (T-EMBLrel. 17, Last sequence update)
DT 01-MAR-2003 (T-EMBLrel. 23, Last annotation update)
DE MHC class I related protein MRL isoform.
GN MRL.
OS Pongo pygmaeus (Orangutan).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pongo.
OC NCBI_TaxID=9600;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=99003494; PubMed=9784382;
RA Yamaguchi H., Kurosawa Y., Hashimoto K.;
RT "Expanded genomic organization of conserved mammalian MHC class I-
RL related genes, human MRL and its murine ortholog.";
RL Biochem. Biophys. Res. Commun. 250:558-564(1998).
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
DR EMBL; AJ271828; CAC28215.1; -.
DR HSSP; Q30201; 1A6Z.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig_c1.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGc1; 1.
DR PROSITE; PS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; Transmembrane.
FT NON_TER 1
SQ SEQUENCE 340 AA; 39375 MW; A893952B78725F17 CRC64;

Query Match 35.1%; Score 531.5; DB 7; Length 340;
Best Local Similarity 39.1%; Pred. No. 1.4e-41;
Matches 106; Conservative 51; Mismatches 111; Indels 3; Gaps 3;

QY 4 RSHSLHYLFWGASEQDGLSLFALGYVDQDLFVFDHESRRVERPPTPVSSRISSQMWL 63
DB 23 RTHSLRYFLRGVSDPIHGVPFISVGVDSPHTTIDYDSVTQKEPRAPWAENLADPHWE 82
QY 64 QLSQSLKGDHMTVDFTWIMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDQQA 123
DB 83 RYTQLLRGQWQMFVELKRLQRHYNHSGSHYQRMIGCELLDGGSTTGFLQAYDQDF 141
QY 124 LEFCPDTLDWRAAEPRAPWTKLEWRHKIRARQNRAYLERDCAQIQQLLELGRGVLDQ 183
DB 142 LIENKDTLSLWADVNDVAHTIKRAWEANQHELOQKNWLEBECIAWLKRFLEYGKDTLQRT 201
QY 184 VPLVKVTHHTVSSVTTLCRALNYYPQNTMKWLKDKQPMDAKEFEPRKDVLPNGDGT 242
DB 202 EPLVRVNRKBTFFGVTALFCAHGFPPEIYNTWMNGEEI-VQEMDYGDLPLPSGDGT 260
QY 243 QGWITLAVPPGEQRVTCQVEHPGLDQPLIV 273

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DB 261 QTWASFLDPQSSNLSYCHVEHGVHMLQV 291

RESULT 8
Q9TK3 PRELIMINARY; PRT; 334 AA.
AC Q9TK3;
DT 01-MAY-2000 (T-EMBLrel. 13, Created)
DT 01-MAY-2000 (T-EMBLrel. 13, Last sequence update)
DT 01-MAR-2003 (T-EMBLrel. 23, Last annotation update)
DE MHC class I-related protein MRL (Fragment).
GN MRL.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OC NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RX TISSUE=Placenta;
RA MEDLINE=99003494; PubMed=9784382;
RA Yamaguchi H., Kurosawa Y., Hashimoto K.;
RT "Expanded genomic organization of conserved mammalian MHC class I-
RL related genes, human MRL and its murine ortholog.";
RL Biochem. Biophys. Res. Commun. 250:558-564(1998).
CC -1- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -1- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
DR EMBL; AF073485; AAC72900.1; -.
DR EMBL; AF073484; AAC72900.1; JOINED.
DR HSSP; Q30201; 1A6Z.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig_c1.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGc1; 1.
DR PROSITE; PS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; Transmembrane.
FT NON_TER 1
SQ SEQUENCE 334 AA; 38586 MW; 4C3E3A8248A39BA4 CRC64;

Query Match 35.0%; Score 530.5; DB 7; Length 334;
Best Local Similarity 39.1%; Pred. No. 1.6e-41;
Matches 106; Conservative 50; Mismatches 112; Indels 3; Gaps 3;

QY 4 RSHSLHYLFWGASEQDGLSLFALGYVDQDLFVFDHESRRVERPPTPVSSRISSQMWL 63
DB 16 RTHSLRYFLRGVSDPIHGVPFISVGVDSPHTTIDYDSVTQKEPRAPWAENLADPHWE 75
QY 64 QLSQSLKGDHMTVDFTWIMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDQQA 123
DB 76 RYTQLLRGQWQMFVELKRLQRHYNHSGSHYQRMIGCELLDGGSTTGFLQAYDQDF 134
QY 124 LEFCPDTLDWRAAEPRAPWTKLEWRHKIRARQNRAYLERDCAQIQQLLELGRGVLDQ 183
DB 135 LIENKDTLSLWADVNDVAHTIKRAWEANQHELOQKNWLEBECIAWLKRFLEYGKDTLQRT 194
QY 184 VPLVKVTHHTVSSVTTLCRALNYYPQNTMKWLKDKQPMDAKEFEPRKDVLPNGDGT 242
DB 195 EPLVRVNRKBTFFGVTALFCAHGFPPEIYNTWMNGEEI-VQSIDYGDILPLPSGDGT 253
QY 243 QGWITLAVPPGEQRVTCQVEHPGLDQPLIV 273
DB 254 QAWASFLDPQSSNLSYCHVEHGVHMLQV 284

RESULT 9
Q9NPL2

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OS Q9NPL2 PRELIMINARY; PRT; 341 AA.
AC Q9NPL2; 01-OCT-2000 (TrEMBLrel. 15, Created)
DT 01-OCT-2000 (TrEMBLrel. 15, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Mrl protein.
GN Mrl.
OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=peripheral blood;
RX MEDLINE=20470599; PubMed=11019920;
RA Parra-Cuadrado J.F., Navarro P., Mirones I., Setien F., Oteo M.,
RA Martinez-Naves E.;
RT "A study on the polymorphism of human MHC class I-related Mrl gene and
RT identification of an Mrl-like pseudogene.";
RL Tissue Antigens 56:170-172(2000).
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
CC EMBL; AJ249778; CAB77667.1; -.
DR HSSP; Q30201; 1A6Z.
DR Genew; HGNC:4975; HLALS.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig cl.
DR InterPro; IPR003006; Ig MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR ProDom; PD000050; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PS00835; IG LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
DR Glycoprotein; Transmembrane.
SQ SEQUENCE 341 AA; 39366 MW; 2990C1F3F0A1CAD9 CRC64;
Query Match 35.0%; Score 530.5; DB 4; Length 341;
Best Local Similarity 39.1%; Pred. No. 1.7e-41;
Matches 106; Conservative 50; Mismatches 112; Indels 3; Gaps 3;
QY 4 RSHSLHYLFMGASEQDLGLSLFALGYVDQLEFVYDHSRRVPRTPWSSRISSQMWL 63
Db 23 RTHSLRYFLRGVSDPIHGVPFISVGVDSPHTYDSVTROKEPRAPWAENLADPHWE 82
QY 64 QLSQSLKGWDHMTVDFTWIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGDGQDA 123
Db 83 RYTQLLRGWQOMFKVELKRLQRHNS-GSHTYQRMIGCELLEDGSTTGLQYAYDQDF 141
QY 124 LEFCPTDLWRAAEPRAPWTKLEWRHKIRARONRAYLERDCPAQLQQLLELGRGVLDQ 183
Db 142 LIENKDTLSLWADVNVHTIKQAEANQHLLYQKNWLEECIAWLKRFLEYKQDTLQRT 201
QY 184 VPLAVKVTHTVT-SSVTLTLCRALNYYPQNTMKLKDQKPMDAKEPEPKDVLPGDGT 242
Db 202 EPLVVRNKRKTFPGVTALFCKAHGFYPPEIYTWKNGEEI-VQIDYGDILPSGDGT 260
QY 243 QGMITLAVPGEQRVTCQVHPLGDLQPLIV 273
Db 261 QAWASIELDPQSSNLSYCHVEHCGVHMLQV 291
RESULT 10
Q95460 PRELIMINARY; PRT; 341 AA.
AC Q95460;
DT 01-FEB-1997 (TrEMBLrel. 02, Created)
DT 01-FEB-1997 (TrEMBLrel. 02, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE Class I histocompatibility antigen-like protein.

OS Q9BCU3 PRELIMINARY; PRT; 341 AA.
AC Q9BCU3;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE MHC class I related protein, Mr1B1 isoform.
GN Mr1.
OS Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
OX NCBI_TaxID=9598;
RN [1]
RP SEQUENCE FROM N.A.
RA Martinez-Naves E.;
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
RN [2]

OS Homo sapiens (Human).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
OX NCBI_TaxID=9606;
RN [1]
RP SEQUENCE FROM N.A.
RC TISSUE=thymus;
RX MEDLINE=95350662; PubMed=7624800;
RA Hashimoto K., Hirai M., Kurosawa Y.;
RT "A gene outside the human MHC related to classical HLA class I
RT genes.";
RL Science 269:693-695(1995).
CC -!- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -!- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
CC EMBL; U22963; AAC50174.1; -.
DR HSSP; Q30201; 1A6Z.
DR Genew; HGNC:4975; HLALS.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003006; Ig MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR ProDom; PD000050; MHC_I; 1.
DR PROSITE; PS00835; IG LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; Transmembrane.
SQ SEQUENCE 341 AA; 39366 MW; 2990C1F3F0A1CAD9 CRC64;
Query Match 35.0%; Score 530.5; DB 7; Length 341;
Best Local Similarity 39.1%; Pred. No. 1.7e-41;
Matches 106; Conservative 50; Mismatches 112; Indels 3; Gaps 3;
QY 4 RSHSLHYLFMGASEQDLGLSLFALGYVDQLEFVYDHSRRVPRTPWSSRISSQMWL 63
Db 23 RTHSLRYFLRGVSDPIHGVPFISVGVDSPHTYDSVTROKEPRAPWAENLADPHWE 82
QY 64 QLSQSLKGWDHMTVDFTWIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGDGQDA 123
Db 83 RYTQLLRGWQOMFKVELKRLQRHNS-GSHTYQRMIGCELLEDGSTTGLQYAYDQDF 141
QY 124 LEFCPTDLWRAAEPRAPWTKLEWRHKIRARONRAYLERDCPAQLQQLLELGRGVLDQ 183
Db 142 LIENKDTLSLWADVNVHTIKQAEANQHLLYQKNWLEECIAWLKRFLEYKQDTLQRT 201
QY 184 VPLAVKVTHTVT-SSVTLTLCRALNYYPQNTMKLKDQKPMDAKEPEPKDVLPGDGT 242
Db 202 EPLVVRNKRKTFPGVTALFCKAHGFYPPEIYTWKNGEEI-VQIDYGDILPSGDGT 260
QY 243 QGMITLAVPGEQRVTCQVHPLGDLQPLIV 273
Db 261 QAWASIELDPQSSNLSYCHVEHCGVHMLQV 291
RESULT 11
Q9BCU3 PRELIMINARY; PRT; 341 AA.
AC Q9BCU3;
DT 01-JUN-2001 (TrEMBLrel. 17, Created)
DT 01-JUN-2001 (TrEMBLrel. 17, Last sequence update)
DT 01-MAR-2003 (TrEMBLrel. 23, Last annotation update)
DE MHC class I related protein, Mr1B1 isoform.
GN Mr1.
OS Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
OX NCBI_TaxID=9598;
RN [1]
RP SEQUENCE FROM N.A.
RA Martinez-Naves E.;
RL Submitted (MAR-2000) to the EMBL/GenBank/DBJ databases.
RN [2]

Db 78 BTRNMGSAQNDVRDLKTLRGYYNQSEAGSHITQIRMYGCDVGDGGLLLRGYDQDAYDGA 137
QY 122 DALEFCCPDITLDWRAAEPRAPWTKLEWHRKIRARQNAYLERDCPAQLQQLLELGRGVLD 181
Db 138 DYIALNEDLRSWTAADTAQAQITKKEWAAV-AEQERSYLEGTCVFWLQKYLEMGKDTLQ 196
QY 182 QOVPPPLVKVTHVTSV-TTLRCALNYPONITMKWLKQKQMDAKFEPKQVLPNGDG 240
Db 197 RAEPFKTHVTRHPSFSDGLVLRWALGFYKPEISLTWQREGQD-QSQDMELVETRPSGDG 255
QY 241 TYQGWITLAVPPGGEQRYTCQVHPGLDQPLIVWE 276
Db 256 TFQWAAVVPVPGGEQSYTCHVQHEGLQEPILTRWD 291

RESULT 14

ID Q30990 PRELIMINARY; PRT; 332 AA.
AC Q30990;
DT 01-NOV-1996 (TREMBlrel. 01, Created)
DT 01-NOV-1996 (TREMBlrel. 01, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
DE Chimpanzee MHC class I Ch1a chain (Fragment).
OS Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
OX NCBI_TaxID=9598;
RN [1]
RP SEQUENCE FROM N.A.
RX MEDLINE=89235215; PubMed=2715640;
RA Parham P., Lawlor D.A., Lomen C.E., Ennis P.D.;
RT "Diversity and diversification of HLA-A,B,C alleles.";
RL J. Immunol. 142:3937-3950(1989).
CC -I- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC -I- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
DR EMBL; M24047; AAA35426.1; -.
DR HSSP; Q95352; 1HHK.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig-cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; Transmembrane.
FT NON_TER 332 332
SQ SEQUENCE 332 AA; 37433 MW; 9AAAS5DF979360 CRC64;

Query Match 33.9%; Score 514; DB 7; Length 332;
Best Local Similarity 40.1%; Pred. No. 5.8e-40;
Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7;

QY 5 SLSHLVFMGASQDGLSLFALGYVDQLFVFYDHE--SRVPRTPWSSRISQW 62
Db 26 SHSMRYFTSVSRPGRGEPFRFAVGYVDDTQFVRFDSDAASQRMPEAPWIEQ-GPEYW 84
QY 63 LOLSQSLKGDHMTFVDFWTIMENHNASK-SHTLQVILGCMEQDNS-TEGYWKYGYDG 120
Db 85 DQETRSKAKHSQTRDVLGTLRGYINQSGDSHTIQTMYGCDVGSGRFLRGYRDAYDG 144
QY 121 QDALEFCPDITLDWRAAEPRAPWTKLEWHRKIRARQNAYLERDCPAQLQQLLELGRGVLD 180
Db 145 KDVIALLNEDLRSWTAADTAQAQITKKEWAAH-AAEQRAYLEGTCTVFWLRRYLENGKETL 203
QY 181 DQOVPLVKVTHH-VTSSVTTTLRCALNYPONITMKWLKQKQMDAKFEPKQVLPNGDG 239

Db 204 QRTDPPKTHHTHPISDHEATLRCWALGFYPAEITLTWQDGED-QTQDTLTVETRPAGD 262
QY 240 GTYQGWITLAVPPGGEQRYTCQVHPGLDQPLIVWE 276
Db 263 GTFQWAAVVPVPGGEQRYTCQVHPGLDQPLIVWE 299
RESULT 15
ID Q9TPL7 PRELIMINARY; PRT; 365 AA.
AC Q9TPL7;
DT 01-MAY-2000 (TREMBlrel. 13, Created)
DT 01-MAY-2000 (TREMBlrel. 13, Last sequence update)
DT 01-MAR-2003 (TREMBlrel. 23, Last annotation update)
DE MHC class I antigen.
GN HLA-A OR PATR-A.
OS Pan troglodytes (Chimpanzee).
OC Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
OC Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Pan.
OX NCBI_TaxID=9598;
RN [1]
RP SEQUENCE FROM N.A.
RX STRAIN=35A-1;
RX MEDLINE=99335357; PubMed=10405321;
RA Matsui M., Machida S., Feinstein S.M., Akatsuka T.;
RT "Molecular analyses of five new chimpanzee MHC class I alleles:
RT Implications for differences between evolutionary mechanisms of HLA-A,
RT -B, and -C loci.";
RL Biochem. Biophys. Res. Commun. 261:46-52(1999).
RN [2]
RP SEQUENCE FROM N.A.
RX TISSUE=Blood;
RX MEDLINE=20322475; PubMed=10866106;
RA de Groot N.G., Otting N., Arguello R., Watkins D.I., Doxiadis G.G.M.,
RA Madrigal J.A., Bontrou R.E.;
RT "Major histocompatibility complex class I diversity in a West African
RT chimpanzee population: implications for HIV research.";
RL Immunogenetics 51:398-409(2000).
CC -I- FUNCTION: INVOLVED IN THE PRESENTATION OF FOREIGN ANTIGENS TO THE
CC IMMUNE SYSTEM (BY SIMILARITY).
CC -I- SUBUNIT: DIMER OF ALPHA CHAIN AND A BETA CHAIN (BETA-2-
CC MICROGLOBULIN) (BY SIMILARITY).
DR EMBL; AF115459; AAF02438.1; -.
DR EMBL; AF168393; AAF72774.1; -.
DR HSSP; Q95352; 1HHK.
DR InterPro; IPR007110; Ig-like.
DR InterPro; IPR003597; Ig-cl.
DR InterPro; IPR003006; Ig_MHC.
DR InterPro; IPR001039; MHC_I.
DR Pfam; PF00047; Ig; 1.
DR Pfam; PF00129; MHC_I; 1.
DR PRINTS; PR01638; MHCCLASSI.
DR ProDom; PD000050; MHC_I; 1.
DR SMART; SM00407; IGcl; 1.
DR PROSITE; PS00835; IG_LIKE; 1.
DR PROSITE; PS00290; IG_MHC; 1.
KW Glycoprotein; MHC; Transmembrane.
SQ SEQUENCE 365 AA; 40819 MW; 4E85F08E33479E38 CRC64;

Query Match 33.9%; Score 514; DB 7; Length 365;
Best Local Similarity 40.1%; Pred. No. 6.5e-40;
Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7;

QY 5 SLSHLVFMGASQDGLSLFALGYVDQLFVFYDHE--SRVPRTPWSSRISQW 62
Db 26 SHSMRYFTSVSRPGRGEPFRFAVGYVDDTQFVRFDSDAASQRMPEAPWIEQ-GPEYW 84
QY 63 LOLSQSLKGDHMTFVDFWTIMENHNASK-SHTLQVILGCMEQDNS-TEGYWKYGYDG 120
Db 85 DQETRSKAKHSQTRDVLGTLRGYINQSGDSHTIQTMYGCDVGSGRFLRGYRDAYDG 144
QY 121 QDALEFCPDITLDWRAAEPRAPWTKLEWHRKIRARQNAYLERDCPAQLQQLLELGRGVLD 180

Db 145 KDYIALNEDLRSWTAADMAAQITKRKWEAAH-AAEQORAYLEGTCTVEWLRRLRYLENGKETL 203
Qy 181 DQOVPLVKVTHH-VTSSVTTLRCALNYYPQNTMKWLKDQKQPMDBAKEPEPKDVLPNGD 239
Db 204 QRTDPPKTHHHPISDHEATLRCWALGFYPAEITLTWQRDGED-QTQDTLVELVETRPAGD 262
Qy 240 GTYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIME 276
Db 263 GTFQKWAAVVPSGEEQRYTCHVQHEGLPKPLTLRWE 299

Search completed: August 5, 2003, 13:10:01
Job time : 33 secs

PF 12-JUN-1998; 98WO-US12436.
 XX
 PR 13-JUN-1997; 97US-0876010.
 XX
 PA (CALY) CALIFORNIA INST OF TECHNOLOGY.
 PA (PROG-) PROGENITOR INC.
 XX
 PI Bjorkman PJ, Feder JN, Schatzman RC;
 XX WPI; 1999-080886/07.
 DR
 XX
 XX New treatment of an iron overload disease - comprises use of HFE
 PT polypeptides provided in a complex with full length, wild type human
 PT (2m), useful in protein replacement therapy
 XX
 XX Claim 5; Page 15; 36pp; English.
 PS
 CC The present sequence represents a H111A/H145A-HFE mutant polypeptide.
 CC The HFE polypeptides (AAW94295-297) provided in a complex with full
 CC length, wild type human beta-2-microglobulin (beta2m) form compositions
 CC in the treatment of primary iron overload diseases (e.g.
 CC hemochromatosis), or other iron overload conditions resulting from
 CC secondary causes (e.g. repeated transfusions). Data regarding the
 CC structure and function correlations of HFE polypeptides is useful in
 CC designing drugs that modulate the HFE gene and HFE activity. The
 CC polypeptides are also useful in protein replacement therapy for
 CC individuals possessing a defective HFE gene (e.g. Hereditary
 CC hemochromatosis). (Antagonists of the polypeptides are also useful in
 CC treating primary and secondary iron overload diseases. The modulators of
 CC the transferrin receptor are useful in treating iron deficiency
 CC conditions such as anemia, and in modulating the amount of iron
 CC transported into a cell. The HFE polypeptides provide a molecular basis
 CC for the relationship between HFE and iron metabolism, which enables
 CC treatment of iron overload and deficiency diseases.
 XX
 SQ Sequence 276 AA;
 Query Match 100.0%; Score 1514; DB 20; Length 276;
 Best Local Similarity 100.0%; Pred. No. 1.1e-132;
 Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 60
 DB 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 60
 QY 61 MWLQLSQSLKGDHMTFTVDFTWMENHNASKESHTLQVILGCEQEDNSTEGYWKYGYDG 120
 DB 61 MWLQLSQSLKGDHMTFTVDFTWMENHNASKESHTLQVILGCEQEDNSTEGYWKYGYDG 120
 QY 121 QDALEFCPDTLDWRAAEPRAMPPTKLEWERHKIRARQNRAVLERDCPAQLQQLLELGRGVL 180
 DB 121 QDALEFCPDTLDWRAAEPRAMPPTKLEWERHKIRARQNRAVLERDCPAQLQQLLELGRGVL 180
 QY 181 DQOVPLVKVTHHTVSSVTLTLCRALNYPQNTMKWLKDKQPMDAKEFEKPDVLPNGDG 240
 DB 181 DQOVPLVKVTHHTVSSVTLTLCRALNYPQNTMKWLKDKQPMDAKEFEKPDVLPNGDG 240
 QY 241 TYCGWITLAVPPGEQRVTCQVEHPGLDQPLIWIWE 276
 DB 241 TYCGWITLAVPPGEQRVTCQVEHPGLDQPLIWIWE 276
 RESULT 2
 ABG72687
 ID ABG72687 standard; protein; 276 AA.
 XX
 AC ABG72687;
 XX
 DT 05-MAR-2003 (first entry)
 XX
 DE Human haemochromatosis (HFE) mature protein, mutant H89A/H123A.
 XX
 XX Human; haemochromatosis; HFE; hereditary haemochromatosis;
 KW

KW iron overload disease; iron deficiency disease; Beta2-microglobulin;
 XX Beta2m; transferrin receptor; anaemia; mutant; mutein.
 OS Homo sapiens.
 OS Synthetic.
 XX
 FH Key Location/Qualifiers
 FT Misc-difference 89 /note= "Wild-type His substituted by Ala"
 FT Misc-difference 123 /note= "Wild-type His substituted by Ala"
 FT
 XX US6391852-B1.
 XX
 XX 21-MAY-2002.
 XX
 XX 12-JUN-1998; 98US-0094964.
 XX
 XX 13-JUN-1997; 97US-0876010.
 XX
 PA (BIRA) BIO-RAD LAB INC.
 PA (CALY) CALIFORNIA INST OF TECHNOLOGY.
 XX
 PI Feder JN, Bjorkman PJ, Schatzman RC;
 XX WPI; 2003-155377/15.
 DR
 XX Method of treating an iron overload disease comprises administration of
 PT a soluble complex comprising a 276 amino acid HFE polypeptide and a
 PT full length, wild-type human beta2m -
 XX
 PS Claim 3; Column 2; 17pp; English.
 XX
 CC The invention relates to a method of treating an iron overload disease
 CC comprising administration of a soluble complex comprising a 276 amino
 CC acid mature HFE (hereditary haemochromatosis gene protein) polypeptide
 CC (ABG72685-ABG72687) and a full length, wild-type human beta2m
 CC (beta2-microglobulin). In a HeLa cell based assay, binding and uptake of
 CC 1,2,51-transferrin in the presence of purified H63D-HFE/beta2m
 CC heterodimers was determined. At a concentration of 250 nM H63D-HFE/
 CC beta2m heterodimers, the transferrin receptor (TfR) displayed a KD for
 CC transferrin of 28 nM. At the same concentration of normal HFE/beta 2m
 CC heterodimers, TfR displayed a KD for transferrin of 40 nM. In the absence
 CC of any HFE/beta 2m heterodimers, TfR displayed a KD for transferrin of
 CC 7nM. It was observed that H63D-HFE/beta 2m heterodimers were 30-40 % less
 CC efficient in decreasing TfR affinity for transferrin compared to
 CC wild-type HFE. The method is useful for treating iron overload diseases
 CC and iron deficiency e.g. anaemia. The present sequence is the H11A/H145A
 CC (residues 111 and 145 of the full length protein, 89/123 of the mature
 CC form) mutant from of mature HFE used to investigate the role of the His
 CC residues in transferrin receptor binding to transferrin.
 XX
 SQ Sequence 276 AA;
 Query Match 100.0%; Score 1514; DB 24; Length 276;
 Best Local Similarity 100.0%; Pred. No. 1.1e-132;
 Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 60
 DB 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 60
 QY 61 MWLQLSQSLKGDHMTFTVDFTWMENHNASKESHTLQVILGCEQEDNSTEGYWKYGYDG 120
 DB 61 MWLQLSQSLKGDHMTFTVDFTWMENHNASKESHTLQVILGCEQEDNSTEGYWKYGYDG 120
 QY 121 QDALEFCPDTLDWRAAEPRAMPPTKLEWERHKIRARQNRAVLERDCPAQLQQLLELGRGVL 180
 DB 121 QDALEFCPDTLDWRAAEPRAMPPTKLEWERHKIRARQNRAVLERDCPAQLQQLLELGRGVL 180
 QY 181 DQOVPLVKVTHHTVSSVTLTLCRALNYPQNTMKWLKDKQPMDAKEFEKPDVLPNGDG 240
 DB 181 DQOVPLVKVTHHTVSSVTLTLCRALNYPQNTMKWLKDKQPMDAKEFEKPDVLPNGDG 240

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QY      241 TYQGWTTLAVPPGEGORYTCQVEHPGLDQPLIVWE 276
DB      241 TYQGWTTLAVPPGEGORYTCQVEHPGLDQPLIVWE 276

RESULT 3
AAW94295
ID      AAW94295 standard; peptide; 276 AA.
XX      AAW94295;
XX      27-APR-1999 (first entry)
XX      Wild-type HFE polypeptide sequence.
DE      HFE; beta-2-microglobulin; beta2m; iron overload; hemochromatosis;
KW      transfusion; protein replacement therapy; hereditary hemochromatosis;
KW      transferrin receptor; iron deficiency; anemia.
XX      Unidentified.
OS      XX
FH      Key Location/Qualifiers
FT      Misc-difference 2 /note= "indicated in the sequence listing as Arg"
PT      XX
XX      W09856814-A1.
XX      17-DEC-1998.
XX      12-JUN-1998; 98WO-US12436.
XX      13-JUN-1997; 97US-0876010.
XX      (CALY ) CALIFORNIA INST OF TECHNOLOGY.
PA      (PROG-) PROGENITOR INC.
XX      Bjorkman PJ, Feder JN, Schatzman RC;
PI      WPI; 1999-080886/07.
XX      New treatment of an iron overload disease - comprises use of HFE
PT      polypeptides provided in a complex with full length, wild type human
PT      (2m), useful in protein replacement therapy
XX      Claim 1; Page 13; 36pp; English.
XX      The present sequence represents a wild-type HFE polypeptide. The HFE
CC      polypeptides (AAW94295-297) provided in a complex with full length,
CC      wild type human beta-2-microglobulin (beta2m) form compositions in the
CC      treatment of primary iron overload diseases (e.g. hemochromatosis), or
CC      other iron overload conditions resulting from secondary causes (e.g.
CC      repeated transfusions). Data regarding the structure and function
CC      correlations of HFE polypeptides is useful in designing drugs that
CC      modulate the HFE gene and HFE activity. The polypeptides are also useful
CC      in protein replacement therapy for individuals possessing a defective
CC      HFE gene (e.g. Hereditary hemochromatosis). (Ant)agonists of the
CC      polypeptides are also useful in treating primary and secondary iron
CC      overload diseases. The modulators of the transferrin receptor are useful
CC      in treating iron deficiency conditions such as anemia, and in modulating
CC      the amount of iron transported into a cell. The HFE polypeptides provide
CC      a molecular basis for the relationship between HFE and iron metabolism,
CC      which enables treatment of iron overload and deficiency diseases.
XX      Sequence 276 AA;
SQ      Query Match 99.2%; Score 1502; DB 20; Length 276;
          Best Local Similarity 99.3%; Pred. No. 1.5e-131;
          Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY      1 RLLRSHLYLFWGASEQDLGLSLFALGVDDQLFVFDHESRRVPRTPWVSSRISQ 60
DB      1 RLLRSHLYLFWGASEQDLGLSLFALGVDDQLFVFDHESRRVPRTPWVSSRISQ 60

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QY      61 MMLQLSLSKGDHMFVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGDG 120
DB      61 MMLQLSLSKGDHMFVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGDG 120

QY      121 QDALEECPTDLDWRAAEPRAPWTKLEWERSHKIRARONRAYLRDPCAOIQQLLELGRGVL 180
DB      121 QDHLSECPDLDWRAAEPRAPWTKLEWERSHKIRARONRAYLRDPCAOIQQLLELGRGVL 180

QY      181 DQQVPLVKVTHVHTSSVTTLCRALNYYYPQNTMKWLKDKQPMDAKEPEPKDVLPGNDG 240
DB      181 DQQVPLVKVTHVHTSSVTTLCRALNYYYPQNTMKWLKDKQPMDAKEPEPKDVLPGNDG 240

QY      241 TYQGWTTLAVPPGEGORYTCQVEHPGLDQPLIVWE 276
DB      241 TYQGWTTLAVPPGEGORYTCQVEHPGLDQPLIVWE 276

RESULT 4
ABG72685
ID      ABG72685 standard; protein; 276 AA.
XX      ABG72685;
XX      05-MAR-2003 (first entry)
XX      Human haemochromatosis (HFE) mature protein.
XX      Human; haemochromatosis; HFE; hereditary haemochromatosis;
KW      iron overload disease; iron deficiency disease; Beta2-microglobulin;
KW      Beta2m; transferrin receptor; anaemia.
XX      Homo sapiens.
XX      US6391852-B1.
XX      21-MAY-2002.
XX      12-JUN-1998; 98US-0094964.
XX      13-JUN-1997; 97US-0876010.
XX      (BIRA ) BIO-RAD LAB INC.
PA      (CALY ) CALIFORNIA INST OF TECHNOLOGY.
XX      Feder JN, Bjorkman PJ, Schatzman RC;
XX      WPI; 2003-155377/15.
XX      Method of treating an iron overload disease comprises administration of
PT      a soluble complex comprising a 276 amino acid HFE polypeptide and a
PT      full length, wild-type human beta2m -
XX      Claim 1; Column 1; 17pp; English.
XX      The invention relates to a method of treating an iron overload disease
CC      comprising administration of a soluble complex comprising a 276 amino
CC      acid mature HFE (hereditary haemochromatosis gene protein) polypeptide
CC      (ABG72685-ABG72687) and a full length, wild-type human beta2m
CC      (beta2-microglobulin). In a HeLa cell based assay, binding and uptake of
CC      125I-transferrin in the presence of purified H63D-HFE/beta2m
CC      heterodimers was determined. At a concentration of 250 nM H63D-HFE/
CC      beta2m heterodimers, the transferrin receptor (TfR) displayed a KD for
CC      transferrin of 28 nM. At the same concentration of normal HFE/beta 2m
CC      heterodimers, TfR displayed a KD for transferrin of 40 nM. In the absence
CC      of any HFE/beta2m heterodimers, TfR displayed a KD for transferrin of
CC      7nM. It was observed that H63D-HFE/beta 2m heterodimers were 30-40 % less
CC      efficient in decreasing TfR affinity for transferrin compared to
CC      wild-type HFE. The method is useful for treating iron overload diseases
CC      and iron deficiency e.g. anaemia. The present sequence is wild-type
CC      mature HFE.
XX      Sequence 276 AA;
SQ

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Query Match 99.2%; Score 1502; DB 24; Length 276;
 Best Local Similarity 99.3%; Pred. No. 1.5e-131;
 Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQDLFVYDHSRRRVEPTPWSSRISSQ 60
 DB 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQDLFVYDHSRRRVEPTPWSSRISSQ 60

QY 61 MWLQSLQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEQEDNSTEGYWKYGYDG 120
 DB 61 MWLQSLQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEQEDNSTEGYWKYGYDG 120

QY 121 QDALEFCPDTLDWRAAPRAWPPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
 DB 121 QDHLEFCPDTLDWRAAPRAWPPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180

QY 181 DQOVPLVKVTHVHTSSVTTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240
 DB 181 DQOVPLVKVTHVHTSSVTTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240

QY 241 TYQGWITLAVPPGEQRVTCQVEHPGLDQPLIVWE 276
 DB 241 TYQGWITLAVPPGEQRVTCQVEHPGLDQPLIVWE 276

RESULT 5

AAW36499

ID AAW36499 standard; Protein; 348 AA.

XX AC

XX AAW36499;

XX DT

XX 14-APR-1998 (first entry)

XX DE

XX Hereditary haemochromatosis gene product.

XX KW

XX Hereditary haemochromatosis; metal toxicity; diagnosis;

XX KW

XX gene therapy; prenatal screening; human.

XX OS

XX Homo sapiens.

XX FH

XX Key Location/Qualifiers

XX FT

XX Misc-difference 63 /note= "substituted by Asp in 24s2 mutant"

XX FT

XX Misc-difference 65 /note= "substituted by Cys in 24d7 variant"

XX FT

XX Misc-difference 282 /note= "substituted by Tyr in 24d1 mutant"

XX FT

XX WO9738137-A1.

XX PD

XX 16-OCT-1997.

XX PF

XX 04-APR-1997; 97WO-US06254.

XX XX

XX 23-MAY-1996; 96US-0652265.

XX PR

XX 04-APR-1996; 96US-0630912.

XX PR

XX 16-APR-1996; 96US-0632673.

XX PA

XX (MERC-) MERCATOR GENETICS INC.

XX XX

XX Drayna DT, Feder JN, Gnirke A, Ruddy D, Thomas WJ;

XX PI

XX Tauchihaashi Z, Wolff RK;

XX XX

XX WPI; 1997-512743/47.

XX DR

XX N-PSDB; AAT96690, AAT96691.

XX XX

XX Hereditary haemochromatosis gene and variants - useful for diagnosis

XX PT

XX and treatment of hereditary haemochromatosis disease

XX XX

XX Disclosure; Fig 4; 115pp; English.

XX PS

XX This polypeptide is the expression product of a novel human gene

XX CC

CC (see AAT96690) whose mutated form is associated with hereditary
 CC haemochromatosis (HH). A single mutation (24d1) in the HH gene
 CC appears responsible for the majority of HH disease. This comprises
 CC a G to A substitution that is present in 86% of affected
 CC chromosomes and in 4% of unaffected chromosomes. It results in a
 CC Cys to Tyr substitution in the encoded protein at a critical
 CC disulphide bridge important for secondary structure. The following
 CC are claimed: the 10825 bp genomic DNA sequence (I), a 1437 bp cDNA
 CC sequence (Ia) (see AAT96691) and their 24d1, 24d2 and 24d7 variants;
 CC a cloning or expression vector; host cells; a peptide product
 CC chosen from the HH gene product, its variants (24d1, 24d2 and
 CC 24d7), or a peptide of at least 56 amino acid residues of these; an
 CC antibody produced using the peptide as an immunogen; a method to
 CC determine the presence or absence of the common HH gene mutation;
 CC an animal model for the HH disease; metal chelation agents, T-cell
 CC differentiation factors and therapeutic agents for the mitigation
 CC of injury due to oxidative process in vivo or mitigation of iron
 CC overload; a method for screening potential therapeutic agents for
 CC activity in connection with HH disease; an antisense oligonucleotide
 CC directed against a transcriptional product of a nucleic acid
 CC sequence as above; and oligonucleotides or pairs of oligonucleotides
 CC covering a range of nucleotides from (I), (Ia) or their variants;
 CC useful for detecting a polymorphism in the HH gene. The invention
 CC also relates to methods for screening for HH homozygotes, to HH
 CC diagnosis, prenatal screening and diagnosis, and therapies of HH
 CC disease, including gene therapy, protein- and antibody-based
 CC therapeutics, and small molecule therapeutics.

XX Sequence 348 AA;

Query Match 99.2%; Score 1502; DB 18; Length 348;
 Best Local Similarity 99.3%; Pred. No. 2e-131;
 Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQDLFVYDHSRRRVEPTPWSSRISSQ 60

DB 23 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQDLFVYDHSRRRVEPTPWSSRISSQ 82

QY 61 MWLQSLQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEQEDNSTEGYWKYGYDG 120

DB 83 MWLQSLQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEQEDNSTEGYWKYGYDG 142

QY 121 QDALEFCPDTLDWRAAPRAWPPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180

DB 143 QDHLEFCPDTLDWRAAPRAWPPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202

QY 181 DQOVPLVKVTHVHTSSVTTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240

DB 203 DQOVPLVKVTHVHTSSVTTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 262

QY 241 TYQGWITLAVPPGEQRVTCQVEHPGLDQPLIVWE 276

DB 263 TYQGWITLAVPPGEQRVTCQVEHPGLDQPLIVWE 298

RESULT 6

AAB19149

ID AAB19149 standard; Protein; 348 AA.

XX AC

XX AAB19149;

XX XX

XX 19-FEB-2001 (first entry)

XX DT

XX A human histocompatibility iron loading (HFE) protein.

XX DE

XX Human; histocompatibility iron loading protein; HFE protein;

XX KW

XX major histocompatibility complex; non-classical class I gene;

XX KW

XX chromosome 6p; iron disorder; haemochromatosis.

XX XX

XX Homo sapiens.

XX OS

XX Key Location/Qualifiers

XX FH

XX Peptide

XX 1..22

FT Misc-difference 63 /note= "signal peptide"
FT FT "when nucleotide 187 is mutated to G, then
FT this residue is Asp"
FT Misc-difference 65 /note= "when nucleotide 193 is mutated to T, then
FT this residue is Cys"
FT Domain 80..108
FT /note= "alpha domain"
FT Misc-difference 93 /note= "when nucleotide 277 is mutated to C, then
FT this residue is Arg"
FT Misc-difference 105 /note= "when nucleotide 314 is mutated to C, then
FT this residue is Thr"
FT FT
PN WO200058515-A1.
XX
XX
XX 05-OCT-2000.
XX
XX 24-MAR-2000; 2000WO-US07982.
XX
XX 26-MAR-1999; 99US-0277457.
XX
XX (BILL-) BILLUPS-ROTHENBERG INC.
XX
XX Rothenberg BE, Sawada-Hirai R, Barton JC;
XX
XX WPI; 2000-647244/62.
DR N-PSDB; AAA96769.
DR
XX
XX Diagnosing an iron disorder e.g. hemochromatosis or a genetic
PT susceptibility to develop it, by determining the presence of a mutation
PT in exon 2 or an intron of a histocompatibility iron loading nucleic
PT acid -
XX
XX Disclosure; Page 3; 55pp; English.
XX
XX The present sequence represents a human histocompatibility iron loading
CC (HFE) protein. The HFE gene is a major histocompatibility (MHC)
CC non-classical class I gene located on chromosome 6p. Mutations in the
CC gene lead to iron disorders. The specification describes a method for
CC diagnosing an iron disorder or a genetic susceptibility to develop of
CC disorder in a mammal. The method comprises determining the presence of
CC a mutation in exon 2 or an intron of a HFE gene or protein. The mutation
CC is not a C to G missense mutation at nucleotide 187 of the sequence
CC given in A96769 (Genbank Accession number U60319). The presence of the
CC mutation indicates the disorder or the genetic susceptibility to the
CC disorder. The method is used to diagnose an iron disorder
CC e.g. haemochromatosis, or a genetic susceptibility to develop it.
XX
XX Sequence 348 AA;
SQ
Query Match 99.2%; Score 1502; DB 21; Length 348;
Best Local Similarity 99.3%; Pred. No. 2e-131;
Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 RLRLSHSLHYLFWGASEQDGLSLFEALGVVDQDLFVFDHESRRVPRTPWVSSRISSQ 60
DB 23 RLRLSHSLHYLFWGASEQDGLSLFEALGVVDQDLFVFDHESRRVPRTPWVSSRISSQ 82
QY 61 MWLQLSLSKLGWDHMTVDFTWIMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSLSKLGWDHMTVDFTWIMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPTDLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCCPAQLQQLLELGRGVL 180
DB 143 QDHLEFCPTDLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCCPAQLQQLLELGRGVL 202
QY 181 DQOVPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
DB 203 DQOVPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262

QY 241 TYQGWITLAVPPGEQRYTCQVEHPGLDQPLIVWE 276
DB 263 TTQGWITLAVPPGEQRYTCQVEHPGLDQPLIVWE 298
RESULT 7
ID AAB36869
XX AAB36869 standard; Protein; 348 AA.
XX
XX AAB36869;
XX
XX 21-FEB-2001 (first entry)
XX Human hereditary hemochromatosis protein.
XX HH; hereditary hemochromatosis; chelation agent;
XX T-cell differentiation factor; iron overload.
XX
XX Homo sapiens.
XX
XX US6140305-A.
XX
XX 31-OCT-2000.
XX
XX 04-APR-1997; 97US-0834497.
XX
XX 04-APR-1996; 96US-0630912.
XX 16-APR-1996; 96US-0632673.
XX 23-MAY-1996; 96US-0652265.
XX
XX (BIRA) BIO-RAD LAB INC.
XX
XX Thomas WJ, Drayna DT, Gnikre A, Ruddy D, Tsuchihashi Z, Wolff RK;
PI Feder JN;
XX
XX WPI; 2001-006341/01.
DR N-PSDB; AAC68425.
XX
XX New hereditary hemochromatosis gene products or polypeptides, useful
PT for treating hereditary hemochromatosis in a patient, and as a metal
PT chelation agent alleviating iron overload -
XX
XX Claim 1; Fig 4; 108pp; English.
XX
XX The present invention relates to hereditary hemochromatosis gene
CC products. These proteins may be used to treat a patient diagnosed as
CC having human hemochromatosis disease. It is also useful as a metal
CC chelation agent or as a T-cell differentiation factor, and for
CC alleviating iron overload. They may also be used in protein replacement
CC therapy for individuals having a defective human hemochromatosis gene.
XX
XX Sequence 348 AA;
SQ
Query Match 99.2%; Score 1502; DB 22; Length 348;
Best Local Similarity 99.3%; Pred. No. 2e-131;
Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
QY 1 RLRLSHSLHYLFWGASEQDGLSLFEALGVVDQDLFVFDHESRRVPRTPWVSSRISSQ 60
DB 23 RLRLSHSLHYLFWGASEQDGLSLFEALGVVDQDLFVFDHESRRVPRTPWVSSRISSQ 82
QY 61 MWLQLSLSKLGWDHMTVDFTWIMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSLSKLGWDHMTVDFTWIMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPTDLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCCPAQLQQLLELGRGVL 180
DB 143 QDHLEFCPTDLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCCPAQLQQLLELGRGVL 202
QY 181 DQOVPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
DB 203 DQOVPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262

QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
 Db 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 298

RESULT 8

AAU80035
 ID AAU80035 standard; Protein; 438 AA.

AC AAU80035;

XX 15-JUL-2002 (first entry)

DE Beta 2 microglobulin (beta2M)/HFE monochain.

XX Human; beta 2 microglobulin; beta2M/HFE monochain; HFE; ischaemia;
 KW iron absorption regulator; intracellular iron absorption; lung injury;
 KW haemochromatosis; transfusion; thalassaemia; haemolytic anaemia;
 KW chronic infection; transferrin receptor; Tfr; brain tumour; cancer;
 KW oxidative stress disorder; tissue damage; vascular disease;
 KW inflammation; atherosclerosis; autoimmune disease;
 KW inflammatory condition.

XX Homo sapiens.

XX WO200224929-A2.

XX 28-MAR-2002.

XX 24-SEP-2001; 2001WO-US29873.

XX 22-SEP-2000; 2000US-234843P.

XX (UYRA-) UNIV RAMOT APPLIED RES & IND DEV LTD.
 PA (MCIN/) MCINNIS P.

XX Ehrlich R, Rotem-Yehudar R, Laham N;

XX WPI; 2002-383192/41.

XX N-PSDB; ABK49917.

XX Soluble beta 2 microglobulin/HFE monochain useful for treating
 PT iron-overload conditions e.g. thalassaemia and chronic infections,
 PT comprises human beta 2 microglobulin linked to alpha domains of HFE by
 PT a linker peptide

XX Example 2; Fig 2; 77pp; English.

XX The invention relates to a soluble polypeptide (I) of beta 2
 CC microglobulin (beta2m)/HFE monochain comprising human beta2m (or its
 CC analogue or active fragment) linked to alpha1-alpha3 domains of human
 CC HFE (a central regulator of iron absorption; undefined), or its analogue
 CC or active fragment, by a flexible linker peptide, or a functional
 CC derivative or salt of (I). (I) is useful for reducing intracellular iron
 CC absorption in patients having hereditary haemochromatosis, transfusions,
 CC thalassaemias, haemolytic anaemia or chronic infections, and for
 CC delivering a therapeutic to cells that over-express transferrin receptor
 CC (Tfr) which are preferably lymphocytes or leukocytes, across the blood-
 CC brain barrier. (I) is further useful for treating brain tumour. (I)
 CC is also useful for treating oxidative stress disorders resulting in
 CC tissue damage e.g. vascular diseases, inflammation, atherosclerosis,
 CC lung injury, ischaemia, etc. A DNA molecule (II) encoding (I) is useful
 CC as a platform for drug delivery of therapeutic use for cancer,
 CC autoimmune diseases and inflammatory conditions. The monochain manifests
 CC specific characteristics advantageous for drug delivery systems. It is a
 CC soluble, stable and fully conformed protein. It binds specifically to
 CC transferrin receptor (Tfr) and therefore targets cells that over-express
 CC this receptor. It is continuously internalised by the target cells, thus
 CC enabling efficient drug delivery. It dissociates from the receptor in the
 CC cells, minimising side effects. It negatively regulates iron absorption,
 CC reducing growth of undesired cells and preventing lymphocyte activation.
 CC It is not diluted in the blood as is transferrin. It should not induce an
 CC immune response since it is a self non-polymorphic protein and delivery of

CC drugs via monochain is expected to overcome drug-resistance since it is a
 CC natural Tfr-binding protein. The present sequence represents the amino
 CC acid sequence of beta2m/HFE monochain.

XX Sequence 438 AA;

Query Match 98.9%; Score 1497; DB 23; Length 438;

Best Local Similarity 99.3%; Pred. No. 7.8e-131;

Matches 273; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYDDQLFVFDHESRRRVEPTPWSSRISSQ 60

Db 135 RLLRSHSLHYLFMGASEQDLGLSLFEALGYDDQLFVFDHESRRRVEPTPWSSRISSQ 194

QY 61 MWLQLSQSLKGWDHMTVDFTIMENHNASKESHTLQVILGCEMEDNSTEGYWKYGYDG 120

Db 195 MWLQLSQSLKGWDHMTVDFTIMENHNASKESHTLQVILGCEMEDNSTEGYWKYGYDG 254

QY 121 QDALEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGYL 180

Db 255 QDHLRFCDPTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGYL 314

QY 181 DQVPPPLVKVTHVTSSVTTLCRALNYPQNTWKWLKDKQPMDAKEPEPKDVLPGD 240

Db 315 DQVPPPLVKVTHVTSSVTTLCRALNYPQNTWKWLKDKQPMDAKEPEPKDVLPGD 374

QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVW 275

Db 375 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVW 409

RESULT 9

AAW94296

ID AAW94296 standard; peptide; 276 AA.

XX AAW94296;

XX 27-APR-1999 (first entry)

XX HFE mutant (H63D-HFE) polypeptide sequence.

XX HFE; beta-2-microglobulin; beta2m; iron overload; hemochromatosis;

XX transfusion; protein replacement therapy; hereditary hemochromatosis;

XX transferrin receptor; iron deficiency; anemia; mutant.

XX Synthetic.

XX Key Location/Qualifiers

FT Misc-difference 2

FT /note= "indicated in the sequence listing as Arg"

FT Misc-difference 41

FT /label= H63D

FT /note= "wild type His (of the mature protein sequence)

FT is replaced by Asp"

PN WO9856814-A1.

XX 17-DEC-1998.

XX 12-JUN-1998; 98WO-US12436.

XX 13-JUN-1997; 97US-0876010.

XX (CALY) CALIFORNIA INST OF TECHNOLOGY.

XX (PROG-) PROGENITOR INC.

XX Bjorkman PJ, Feder JN, Schatzman RC;

XX WPI; 1999-080886/07.

XX New treatment of an iron overload disease - comprises use of HFE

XX polypeptides provided in a complex with full length, wild type human

XX (2m), useful in protein replacement therapy

XX PS Claim 3; Page 14; 36pp; English.

XX CC The present sequence represents a H63D-HFE mutant polypeptide. The HFE

XX CC polypeptides (AAW94295-297) provided in a complex with full length,

XX CC wild type human beta-2-microglobulin (beta2m) form compositions in the

XX CC treatment of primary iron overload diseases (e.g. hemochromatosis), or

XX CC other iron overload conditions resulting from secondary causes (e.g.

XX CC repeated transfusions). Data regarding the structure and function

XX CC correlate of HFE polypeptides is useful in designing drugs that

XX CC modulate the HFE gene and HFE activity. The polypeptides are also useful

XX CC in protein replacement therapy for individuals possessing a defective

XX CC HFE gene (e.g. Hereditary hemochromatosis). (Antagonists of the

XX CC polypeptides are also useful in treating primary and secondary iron

XX CC overload diseases. The modulators of the transferrin receptor are useful

XX CC in treating iron deficiency conditions such as anemia, and in modulating

XX CC the amount of iron transported into a cell. The HFE polypeptides provide

XX CC a molecular basis for the relationship between HFE and iron metabolism,

XX CC which enables treatment of iron overload and deficiency diseases.

XX SQ Sequence 276 AA;

Query Match 98.6%; Score 1493; DB 20; Length 276;

Best Local Similarity 98.9%; Pred. No. 1e-130;

Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISSQ 60

DB 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISSQ 60

QY 61 MWLQSLQSLKGDHMTFTVDFWTIMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 120

DB 61 MWLQSLQSLKGDHMTFTVDFWTIMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 120

QY 121 QDALEFCFDPDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180

DB 121 QDHLEFCFDPDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180

QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNITMKWLKDKQPMDAKEPEPKDVLPGDNG 240

DB 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNITMKWLKDKQPMDAKEPEPKDVLPGDNG 240

QY 241 TYQGWITLAVPPGEQRVTCQVEHPGLDQPLIWIWE 276

DB 241 TYQGWITLAVPPGEQRVTCQVEHPGLDQPLIWIWE 276

RESULT 10

ABG72686

ID ABG72686 standard; protein; 276 AA.

XX AC ABG72686;

XX DT 05-MAR-2003 (first entry)

XX DE Human haemochromatosis (HFE) mature protein, mutant H41D.

XX KW Human; haemochromatosis; HFE; hereditary haemochromatosis;

XX KW iron overload disease; iron deficiency disease; Beta2-microglobulin;

XX KW Beta2m; transferrin receptor; anaemia; mutant; mutein.

XX OS Homo sapiens.

XX OS Synthetic.

XX FH Key

XX FT Location/Qualifiers

XX FT Misc-difference 41

XX FT /note= "Wild-type His substituted by Asp"

XX PN US6391852-B1.

XX PD 21-MAY-2002.

XX PF 12-JUN-1998; 98US-0094964.

XX PR 13-JUN-1997; 97US-0876010.

XX PA (BIRA) BIO-RAD LAB INC.

XX PA (CALY) CALIFORNIA INST OF TECHNOLOGY.

XX PI Feder JN, Bjorkman PU, Schatzman RC;

XX XX WPI; 2003-155377/15.

XX PT Method of treating an iron overload disease comprises administration of

XX PT a soluble complex comprising a 276 amino acid HFE polypeptide and a

XX PT full length, wild-type human beta2m -

XX PS Claim 2; Column 2; 17pp; English.

XX CC The invention relates to a method of treating an iron overload disease

XX CC comprising administration of a soluble complex comprising a 276 amino

XX CC acid mature HFE (hereditary haemochromatosis gene protein) polypeptide

XX CC (ABG72685-ABG72687) and a full length, wild-type human beta2m

XX CC (beta2-microglobulin). In a HeLa cell based assay, binding and uptake of

XX CC ⁵¹Fe-transferrin in the presence of purified H63D-HFE/beta2m

XX CC heterodimers was determined. At a concentration of 250 nM H63D-HFE/

XX CC beta2m heterodimers, the transferrin receptor (TfR) displayed a KD for

XX CC transferrin of 28 nM. At the same concentration of normal HFE/beta2m

XX CC heterodimers, TfR displayed a KD for transferrin of 40 nM. In the absence

XX CC of any HFE/beta2m heterodimers, TfR displayed a KD for transferrin of

XX CC 7nM. It was observed that H63D-HFE/beta2m heterodimers were 30-40 % less

XX CC efficient in decreasing TfR affinity for transferrin compared to

XX CC wild-type HFE. The method is useful for treating iron overload diseases

XX CC and iron deficiency e.g. anaemia. The present sequence is the H63D

XX CC (residue 63 of the full length protein, 41 of the mature form)

XX CC mutant form of mature HFE used to investigate the role of the His

XX CC residue in transferrin receptor binding to transferrin.

XX SQ Sequence 276 AA;

Query Match 98.6%; Score 1493; DB 24; Length 276;

Best Local Similarity 98.9%; Pred. No. 1e-130;

Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISSQ 60

DB 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISSQ 60

QY 61 MWLQSLQSLKGDHMTFTVDFWTIMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 120

DB 61 MWLQSLQSLKGDHMTFTVDFWTIMENHNASKESHTLQVILGCMQEDNSTEGYWKYGYDG 120

QY 121 QDALEFCFDPDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180

DB 121 QDHLEFCFDPDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180

QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNITMKWLKDKQPMDAKEPEPKDVLPGDNG 240

DB 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNITMKWLKDKQPMDAKEPEPKDVLPGDNG 240

QY 241 TYQGWITLAVPPGEQRVTCQVEHPGLDQPLIWIWE 276

DB 241 TYQGWITLAVPPGEQRVTCQVEHPGLDQPLIWIWE 276

RESULT 11

AAB36871

ID AAB36871 standard; Protein; 348 AA.

XX AC AAB36871;

XX DT 21-FEB-2001 (first entry)

XX DE Human hereditary hemochromatosis 24d2 mutation protein.

XX XX HH; hereditary hemochromatosis; chelation agent;

KW T-cell differentiation factor; iron overload.

XX Homo sapiens.

OS US6140305-A.

PN 31-OCT-2000.

XX 04-APR-1997; 97US-0834497.

XX 04-APR-1996; 96US-0630912.

PR 16-APR-1996; 96US-0632673.

PR 23-MAY-1996; 96US-0652265.

XX (BIRA) BIO-RAD LAB INC.

XX Thomas WJ, Drayna DT, Gnirke A, Ruddy D, Tsuchihashi Z, Wolff RK;

PI Feder JN;

DR WPI; 2001-006341/01.

DR N-PSDB; AAC68427.

XX New hereditary hemochromatosis gene products or polypeptides, useful

PT for treating hereditary hemochromatosis in a patient, and as a metal

PT chelation agent alleviating iron overload -

XX Claim 3; Fig 4; 108pp; English.

XX The present invention relates to hereditary hemochromatosis gene

CC products. These proteins may be used to treat a patient diagnosed as

CC having human hemochromatosis disease. It is also useful as a metal

CC chelation agent or as a T-cell differentiation factor, and for

CC alleviating iron overload. They may also be used in protein replacement

CC therapy for individuals having a defective human hemochromatosis gene.

XX SQ Sequence 348 AA;

Query Match 98.6%; Score 1493; DB 22; Length 348;

Best Local Similarity 98.9%; Pred. No. 1.4e-130;

Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 60

DB 23 RLLRSHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 82

QY 61 MWLQLSQSLKGDHMFVDFWTIMENHNHKSHTLQVILGCEMOEDNSTEGYWKYGYDG 120

DB 83 MWLQLSQSLKGDHMFVDFWTIMENHNHKSHTLQVILGCEMOEDNSTEGYWKYGYDG 142

QY 121 QDALEFCPTDLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180

DB 143 QDHLEFCPTDLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 202

QY 181 DQVPPPLVKVTHVTSSVTLRCLALNYYPQNTMKWLKDKQPMDAKEFEPKDVLPNGDG 240

DB 203 DQVPPPLVKVTHVTSSVTLRCLALNYYPQNTMKWLKDKQPMDAKEFEPKDVLPNGDG 262

QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276

DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 298

RESULT 12

AAB36870

ID AAB36870 standard; Protein; 348 AA.

XX AAB36870;

XX 21-FEB-2001 (first entry)

XX Human hereditary hemochromatosis 24d1 mutation protein.

XX HH; hereditary hemochromatosis; chelation agent;

KW

KW T-cell differentiation factor; iron overload.

XX Homo sapiens.

OS US6140305-A.

PN 31-OCT-2000.

XX 04-APR-1997; 97US-0834497.

XX 04-APR-1996; 96US-0630912.

PR 16-APR-1996; 96US-0632673.

PR 23-MAY-1996; 96US-0652265.

XX (BIRA) BIO-RAD LAB INC.

XX Thomas WJ, Drayna DT, Gnirke A, Ruddy D, Tsuchihashi Z, Wolff RK;

PI Feder JN;

DR WPI; 2001-006341/01.

DR N-PSDB; AAC68426.

XX New hereditary hemochromatosis gene products or polypeptides, useful

PT for treating hereditary hemochromatosis in a patient, and as a metal

PT chelation agent alleviating iron overload -

XX Claim 2; Fig 3; 108pp; English.

XX The present invention relates to hereditary hemochromatosis gene

CC products. These proteins may be used to treat a patient diagnosed as

CC having human hemochromatosis disease. It is also useful as a metal

CC chelation agent or as a T-cell differentiation factor, and for

CC alleviating iron overload. They may also be used in protein replacement

CC therapy for individuals having a defective human hemochromatosis gene.

XX SQ Sequence 348 AA;

Query Match 98.5%; Score 1491; DB 22; Length 348;

Best Local Similarity 98.9%; Pred. No. 2.1e-130;

Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 60

DB 23 RLLRSHSLHYLFMGASQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISSQ 82

QY 61 MWLQLSQSLKGDHMFVDFWTIMENHNHKSHTLQVILGCEMOEDNSTEGYWKYGYDG 120

DB 83 MWLQLSQSLKGDHMFVDFWTIMENHNHKSHTLQVILGCEMOEDNSTEGYWKYGYDG 142

QY 121 QDALEFCPTDLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180

DB 143 QDHLEFCPTDLDWRAAEPRAPWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 202

QY 181 DQVPPPLVKVTHVTSSVTLRCLALNYYPQNTMKWLKDKQPMDAKEFEPKDVLPNGDG 240

DB 203 DQVPPPLVKVTHVTSSVTLRCLALNYYPQNTMKWLKDKQPMDAKEFEPKDVLPNGDG 262

QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276

DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 298

RESULT 13

AAB36872

ID AAB36872 standard; Protein; 348 AA.

XX AAB36872;

XX 21-FEB-2001 (first entry)

XX Human hereditary hemochromatosis 24d1/2 mutation protein.

XX HH; hereditary hemochromatosis; chelation agent;

KW

```
KW T-cell differentiation factor; iron overload.
XX Homo sapiens.
XX US6140305-A.
XX 31-OCT-2000.
XX
XX 04-APR-1997; 97US-0834497.
XX
XX 04-APR-1996; 96US-0630912.
XX 16-APR-1996; 96US-0632673.
XX 23-MAY-1996; 96US-0652265.
XX
XX (BIRA ) BIO-RAD LAB INC.
XX
XX Thomas WJ, Drayna DT, Gnirke A, Ruddy D, Tsuchihashi Z, Wolff RK;
XX Feder JN;
XX
XX WPI; 2001-006341/01.
XX N-PSDB; AAC68428.
XX
XX New hereditary hemochromatosis gene products or polypeptides, useful
XX PT for treating hereditary hemochromatosis in a patient, and as a metal
XX PT chelation agent alleviating iron overload -
XX
XX Claim 4; Fig 4; 108pp; English.
XX
XX The present invention relates to hereditary hemochromatosis gene
XX products. These proteins may be used to treat a patient diagnosed as
XX having human hemochromatosis disease. It is also useful as a metal
XX chelation agent or as a T-cell differentiation factor, and for
XX CC alleviating iron overload. They may also be used in protein replacement
XX CC therapy for individuals having a defective human hemochromatosis gene.
XX
XX Sequence 348 AA;
SQ
Query Match 97.9%; Score 1482; DB 22; Length 348;
Best Local Similarity 98.6%; Pred. No. 1.5e-129;
Matches 272; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
QY 1 RLLRSHLVLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWYSSRISSQ 60
DB 23 RLLRSHLVLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWYSSRISSQ 82
QY 61 MWLQSLQSLKGWDMFTVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGWDMFTVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQOVPLVKVTHVTSSVTTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
DB 203 DQOVPLVKVTHVTSSVTTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262
QY 241 TYQGWITLAVPPGEORVTCOVHPGLDQPLIWIWE 276
DB 263 TYQGWITLAVPPGEORVTCOVHPGLDQPLIWIWE 298
RESULT 14
AAB36873
ID AAB36873 standard; Protein; 361 AA.
XX
XX AAB36873;
XX
XX 21-FEB-2001 (first entry)
XX
XX Rabbit leukocyte antigen.
XX
XX HH; hereditary hemochromatosis; chelation agent;
KW
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KW T-cell differentiation factor; iron overload.
XX Oryctolagus cuniculus.
XX US6140305-A.
XX 31-OCT-2000.
XX
XX 04-APR-1997; 97US-0834497.
XX
XX 04-APR-1996; 96US-0630912.
XX 16-APR-1996; 96US-0632673.
XX 23-MAY-1996; 96US-0652265.
XX
XX (BIRA ) BIO-RAD LAB INC.
XX
XX Thomas WJ, Drayna DT, Gnirke A, Ruddy D, Tsuchihashi Z, Wolff RK;
XX Feder JN;
XX
XX WPI; 2001-006341/01.
XX
XX New hereditary hemochromatosis gene products or polypeptides, useful
XX PT for treating hereditary hemochromatosis in a patient, and as a metal
XX PT chelation agent alleviating iron overload -
XX
XX Disclosure; Fig 7; 108pp; English.
XX
XX The present invention relates to hereditary hemochromatosis gene
XX products. These proteins may be used to treat a patient diagnosed as
XX having human hemochromatosis disease. It is also useful as a metal
XX chelation agent or as a T-cell differentiation factor, and for
XX CC alleviating iron overload. They may also be used in protein replacement
XX CC therapy for individuals having a defective human hemochromatosis gene.
XX
XX Sequence 361 AA;
SQ
Query Match 34.1%; Score 517; DB 22; Length 361;
Best Local Similarity 40.1%; Pred. No. 1.1e-39;
Matches 111; Conservative 43; Mismatches 115; Indels 8; Gaps 7;
QY 5 SLSLHVLFMGASEQDGLSLFEALGYVDDQLFVFDHE--SRVPRTPWYSSRISSQMW 62
DB 26 SLSLHVLFMGASEQDGLSLFEALGYVDDQLFVFDHE--SRVPRTPWYSSRISSQMW 84
QY 63 LQLSLSLQSLKGWDMFTVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 85 DQQTQIAKDTATQTFRVNLTALRYNQSAAGSHITQTFGCEVWADGRFFHGYRQYAYDG 144
QY 121 QDALEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 145 ADVIALNEDLRGWSAATAAQNTQKWEAAG-EAERHAYLERECVEMLRRLYLENGKETL 203
QY 181 DQOVPLVKVTHVTSS-VTTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 239
DB 204 QDALEFCPDTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 262
QY 240 TYQGWITLAVPPGEORVTCOVHPGLDQPLIWIWE 276
DB 263 TYQGWITLAVPPGEORVTCOVHPGLDQPLIWIWE 299
RESULT 15
AAB36874
ID AAB36874 standard; Protein; 365 AA.
XX
XX AAB36874;
XX
XX 21-FEB-2001 (first entry)
XX
XX MHC class I protein.
XX
XX HH; hereditary hemochromatosis; chelation agent;
KW T-cell differentiation factor; iron overload.
KW
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GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:10:04 ; Search time 33 Seconds
(without alignments)
993.264 Million cell updates/sec

Title: US-10-092-404-3
Perfect score: 1514
Sequence: 1 RLRLSHLHFLMGASEQDL.....RYTCQVHPGLDQPLIVWE 276

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 451899 seqs, 118759770 residues

Total number of hits satisfying chosen parameters: 451899

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Published Applications AA:
1: /cgn2_6/ptodata/1/pubpaa/US07_PUBCOMB.pep.*
2: /cgn2_6/ptodata/1/pubpaa/PCT_NEW_PUB.pep.*
3: /cgn2_6/ptodata/1/pubpaa/US06_NEW_PUB.pep.*
4: /cgn2_6/ptodata/1/pubpaa/US06_PUBCOMB.pep.*
5: /cgn2_6/ptodata/1/pubpaa/US07_NEW_PUB.pep.*
6: /cgn2_6/ptodata/1/pubpaa/PCTUS_PUBCOMB.pep.*
7: /cgn2_6/ptodata/1/pubpaa/US08_NEW_PUB.pep.*
8: /cgn2_6/ptodata/1/pubpaa/US08_PUBCOMB.pep.*
9: /cgn2_6/ptodata/1/pubpaa/US09A_PUBCOMB.pep.*
10: /cgn2_6/ptodata/1/pubpaa/US09B_PUBCOMB.pep.*
11: /cgn2_6/ptodata/1/pubpaa/US09C_PUBCOMB.pep.*
12: /cgn2_6/ptodata/1/pubpaa/US09_NEW_PUB.pep.*
13: /cgn2_6/ptodata/1/pubpaa/US10A_PUBCOMB.pep.*
14: /cgn2_6/ptodata/1/pubpaa/US10B_PUBCOMB.pep.*
15: /cgn2_6/ptodata/1/pubpaa/US10C_PUBCOMB.pep.*
16: /cgn2_6/ptodata/1/pubpaa/US10_NEW_PUB.pep.*
17: /cgn2_6/ptodata/1/pubpaa/US60_NEW_PUB.pep.*
18: /cgn2_6/ptodata/1/pubpaa/US60_PUBCOMB.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1514	100.0	276	15	US-10-092-404-3
2	1502	99.2	276	15	US-10-092-404-1
3	1502	99.2	348	12	US-09-981-606-2
4	1493	98.6	276	15	US-10-092-404-2
5	504	33.3	92	14	US-10-016-634A-120
6	500	33.0	280	15	US-10-073-300-6
7	500	33.0	415	15	US-10-073-300-5
8	486	32.1	298	15	US-10-205-823-40
9	486	32.1	298	15	US-10-205-823-42
10	486	32.1	298	15	US-10-177-293-23
11	471	31.1	542	15	US-10-015-535-32
12	471	31.1	542	15	US-10-015-535-34
13	470	31.0	542	15	US-10-015-535-36
14	468	30.9	540	15	US-10-015-535-22
15	468	30.9	541	15	US-10-015-535-28

16	468	30.9	542	15	US-10-015-535-24	Sequence 24, Appl
17	468	30.9	542	15	US-10-015-535-26	Sequence 26, Appl
18	446	29.5	332	9	US-09-870-521-3	Sequence 3, Appl
19	441	29.1	334	9	US-09-870-521-4	Sequence 4, Appl
20	439	29.0	540	15	US-10-015-535-30	Sequence 30, Appl
21	354.5	23.4	170	9	US-09-925-301-1307	Sequence 1307, Appl
22	330	21.8	271	9	US-09-925-301-1431	Sequence 1431, Ap
23	275	18.2	145	9	US-09-810-560-8	Sequence 8, Appl
24	273	18.0	181	11	US-09-013-077A-13	Sequence 13, Appl
25	237	15.7	184	10	US-09-858-580-21	Sequence 21, Appl
26	237	15.7	184	11	US-09-847-172-21	Sequence 21, Appl
27	226	14.9	91	9	US-09-864-761-38005	Sequence 38005, A
28	223	14.7	91	9	US-09-864-761-35461	Sequence 35461, A
29	217.5	14.4	171	15	US-10-144-929-1116	Sequence 1116, App
30	210.5	13.9	104	9	US-09-925-302-835	Sequence 835, App
31	207	13.7	117	9	US-09-810-560-9	Sequence 9, Appl
32	202.5	13.4	183	15	US-10-036-542-62	Sequence 62, Appl
33	196.5	13.0	93	9	US-09-864-761-39479	Sequence 39479, A
34	196.5	13.0	110	9	US-09-864-761-35339	Sequence 35339, A
35	196.5	13.0	114	9	US-09-864-761-37988	Sequence 37988, A
36	176.5	11.7	261	10	US-09-925-664-30	Sequence 30, Appl
37	173	11.4	110	10	US-09-796-692-799	Sequence 799, App
38	173	11.4	110	10	US-09-796-692-2139	Sequence 2139, Ap
39	173	11.4	110	15	US-10-040-862-799	Sequence 799, App
40	173	11.4	110	15	US-10-040-862-2139	Sequence 2139, Ap
41	170	11.2	411	14	US-10-015-536-17	Sequence 17, Appl
42	167.5	11.1	285	10	US-09-756-983-24	Sequence 24, Appl
43	167	11.0	246	9	US-09-989-722-225	Sequence 225, App
44	167	11.0	246	9	US-09-989-723-225	Sequence 225, App
45	167	11.0	246	9	US-09-989-279-225	Sequence 225, App

ALIGNMENTS

RESULT 1

US-10-092-404-3

; Sequence 3, Application US/10092404

; Publication No. US20030073627A1

; GENERAL INFORMATION:

; APPLICANT: Feder, John N.

; Bjorkman, Pamela J.

; Schatzman, Randall C.

; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR

; DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES

; AND IRON DEFICIENCY DISEASES

; NUMBER OF SEQUENCES: 5

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Pennie & Edmonds, LLP

; STREET: 1155 Avenue of the Americas

; CITY: New York

; STATE: NY

; COUNTRY: USA

; ZIP: 10036-2811

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Diskette

; OPERATING SYSTEM: Windows

; SOFTWARE: FASTSEQ for Windows Version 2.0b

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/10/092,404

; FILING DATE: 04-Mar-2002

; CLASSIFICATION: <Unknown>

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: US/09/094,964

; FILING DATE: June 12, 1998

; APPLICATION NUMBER: 08/876,010

; FILING DATE: June 13, 1997

; ATTORNEY/AGENT INFORMATION:

; NAME: Poissant, Brian M

; REGISTRATION NUMBER: 28,462

; REFERENCE/DOCKET NUMBER: 8907-0074-999

; TELECOMMUNICATION INFORMATION:

TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 3:
SEQUENCE CHARACTERISTICS:
LENGTH: 276 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
SEQUENCE DESCRIPTION: SEQ ID NO: 3:
US-10-092-404-3

Query Match 100.0%; Score 1514; DB 15; Length 276;
Best Local Similarity 100.0%; Pred. No. 1.8e-147;
Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWVSSRISSQ 60
DB 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWVSSRISSQ 60
QY 61 MWLQSLKSGWDMFTVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 61 MWLQSLKSGWDMFTVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
QY 121 QDALEFCPTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCAQLOLLELGRGVL 180
DB 121 QDALEFCPTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCAQLOLLELGRGVL 180
QY 181 DQVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
DB 181 DQVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 276
DB 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 276

RESULT 2

US-10-092-404-1
Sequence 1, Application US/10092404
Publication No. US20030073627A1
GENERAL INFORMATION:
APPLICANT: Feder, John N.
Bjorkman, Pamela J.
Schatzman, Randall C.
TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
AND IRON DEFICIENCY DISEASES
NUMBER OF SEQUENCES: 5
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds, LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: NY
COUNTRY: USA
ZIP: 10036-2811
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: Windows
SOFTWARE: FastSeq for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/10/092,404
FILING DATE: 04-Mar-2002
CLASSIFICATION: <Unknown>
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US/09/094,964
FILING DATE: June 12, 1998
APPLICATION NUMBER: 08/876,010
FILING DATE: June 13, 1997
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M

REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0074-999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 276 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
SEQUENCE DESCRIPTION: SEQ ID NO: 1:
US-10-092-404-1

Query Match 99.2%; Score 1502; DB 15; Length 276;
Best Local Similarity 99.3%; Pred. No. 3.1e-146;
Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWVSSRISSQ 60
DB 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWVSSRISSQ 60
QY 61 MWLQSLKSGWDMFTVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 61 MWLQSLKSGWDMFTVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
QY 121 QDALEFCPTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCAQLOLLELGRGVL 180
DB 121 QDALEFCPTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCAQLOLLELGRGVL 180
QY 181 DQVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
DB 181 DQVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 276
DB 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 276

RESULT 3

US-09-981-606-2
Sequence 2, Application US/09981606
Publication No. US20030129595A1
GENERAL INFORMATION:
APPLICANT: Rothenberg et al.
TITLE OF INVENTION: Mutations associated with iron disorders
FILE REFERENCE: 24065-004CCN
CURRENT APPLICATION NUMBER: US/09/981,606
CURRENT FILING DATE: 2002-10-16
PRIOR APPLICATION NUMBER: 09/277,457
PRIOR FILING DATE: 1999-03-26
NUMBER OF SEQ ID NOS: 30
SOFTWARE: PatentIn Ver. 2.1
SEQ ID NO 2
LENGTH: 348
TYPE: PRT
ORGANISM: Homo sapiens
US-09-981-606-2

Query Match 99.2%; Score 1502; DB 12; Length 348;
Best Local Similarity 99.3%; Pred. No. 4.3e-146;
Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWVSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWVSSRISSQ 82
QY 61 MWLQSLKSGWDMFTVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLKSGWDMFTVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142

QY 121 QDALEFCPTDLWRAAEPRAPWTKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGVL 180
Db 143 QDHFECPTDLWRAAEPRAPWTKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGVL 202
QY 181 DQOVPLVKVTHVTSSVTTLCRALNYPQNTMKWLKDKQPMDAKEFEKPDVLPNGDG 240
Db 203 DQOVPLVKVTHVTSSVTTLCRALNYPQNTMKWLKDKQPMDAKEFEKPDVLPNGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
Db 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 298

RESULT 4
US-10-092-404-2
; Sequence 2, Application US/10092404
; Publication No. US20030073627A1
; GENERAL INFORMATION:
; APPLICANT: Feder, John N.
; Bjorkman, Pamela J.
; Schatzman, Randall C.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
; DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
; AND IRON DEFICIENCY DISEASES
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds, LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10036-2811
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: Windows
; SOFTWARE: FastSeq for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/10/092,404
; FILING DATE: 04-Mar-2002
; CLASSIFICATION: <Unknown>
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/09/094,964
; FILING DATE: June 12, 1998
; APPLICATION NUMBER: 08/876,010
; FILING DATE: June 13, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0074-999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-493-4935
; TELEFAX: 650-493-5556
; TELEX: 66141 PENNIE
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 276 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; SEQUENCE DESCRIPTION: SEQ ID NO: 2:
US-10-092-404-2

Query Match 98.6%; Score 1493; DB 15; Length 276;
Best Local Similarity 98.9%; Pred. No. 2.7e-145;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLLRSHSLYLFMGASEQDLGLSLFALGYVDDQLFVFDHESRRYPRTPWVSSRISQ 60
Db 1 RLLRSHSLYLFMGASEQDLGLSLFALGYVDDQLFVFDHESRRYPRTPWVSSRISQ 60
QY 61 MMLQLSLSLKGNDHMTFVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120

Db 61 MMLQLSLSLKGNDHMTFVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
QY 121 QDALEFCPTDLWRAAEPRAPWTKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGVL 180
Db 121 QDHFECPTDLWRAAEPRAPWTKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGVL 180
QY 181 DQOVPLVKVTHVTSSVTTLCRALNYPQNTMKWLKDKQPMDAKEFEKPDVLPNGDG 240
Db 181 DQOVPLVKVTHVTSSVTTLCRALNYPQNTMKWLKDKQPMDAKEFEKPDVLPNGDG 240
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
Db 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276

RESULT 5

US-10-016-634A-120
; Sequence 120, Application US/10016634A
; Publication No. US20020192666A1
; GENERAL INFORMATION:
; APPLICANT: Sun, Yongming
; APPLICANT: Recipon, Herve
; APPLICANT: Ghosh, Malavika
; APPLICANT: Liu, Changhai
; TITLE OF INVENTION: Compositions and Methods Relating to Colon Specific Genes and Pri
; FILE REFERENCE: DEX-0255
; CURRENT APPLICATION NUMBER: US/10/016,634A
; CURRENT FILING DATE: 2001-10-31
; PRIOR APPLICATION NUMBER: US 60/244,258
; PRIOR FILING DATE: 2000-10-31
; NUMBER OF SEQ ID NOS: 176
; SOFTWARE: PatentIn version 3.1
; SEQ ID NO 120
; LENGTH: 92
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-016-634A-120

Query Match 33.3%; Score 504; DB 14; Length 92;
Best Local Similarity 98.9%; Pred. No. 3.1e-44;
Matches 91; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 92 ESHTLQVILGCEMQEDNSTEGYWKYGYDGQDALEFCPTDLWRAAEPRAPWTKLEWERHK 151
Db 1 ESHTLQVILGCEMQEDNSTEGYWKYGYDGQDHFECPTDLWRAAEPRAPWTKLEWERHK 60
QY 152 IRARONRAYLERDPCPAQLQQLLELGRGVLDDQ 183
Db 61 IRARONRAYLERDPCPAQLQQLLELGRGVLDDQ 92

RESULT 6

US-10-073-300-6
; Sequence 6, Application US/10073300
; Publication No. US20030003535A1
; GENERAL INFORMATION:
; APPLICANT: Reiter, Yoram
; TITLE OF INVENTION: SINGLE CHAIN CLASS I MAJOR HISTO- COMPATIBILITY COMPLEXES
; FILE REFERENCE: 02/23339
; CURRENT APPLICATION NUMBER: US/10/073,300
; CURRENT FILING DATE: 2002-06-25
; NUMBER OF SEQ ID NOS: 20
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 6
; LENGTH: 280
; TYPE: PRT
; ORGANISM: Homo sapiens
US-10-073-300-6

Query Match 33.0%; Score 500; DB 15; Length 280;
Best Local Similarity 39.4%; Pred. No. 3.6e-43;
Matches 109; Conservative 44; Mismatches 116; Indels 8; Gaps 7;

APPLICANT: Zhao, Xumei
TITLE OF INVENTION: NOVEL GENES, COMPOSITIONS, KITS, AND
METHODS FOR IDENTIFICATION, ASSESSMENT, PREVENTION, AND
THERAPY OF PROSTATE CANCER
FILE REFERENCE: MRI-044
CURRENT APPLICATION NUMBER: US/10/205,823
CURRENT FILING DATE: 2002-07-25
PRIOR APPLICATION NUMBER: 60/307,992
PRIOR FILING DATE: 2001-07-25
PRIOR APPLICATION NUMBER: 60/314,356
PRIOR FILING DATE: 2001-08-22
PRIOR APPLICATION NUMBER: 60/325,020
PRIOR FILING DATE: 2001-09-25
PRIOR APPLICATION NUMBER: 60/341,746
PRIOR FILING DATE: 2001-12-12
PRIOR APPLICATION NUMBER: 60/362,158
PRIOR FILING DATE: 2002-03-05
NUMBER OF SEQ ID NOS: 455
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 42
LENGTH: 298
TYPE: PRT
ORGANISM: Homo sapiens
US-10-205-823-42

Query Match 32.1%; Score 486; DB 15; Length 298;
Best Local Similarity 36.7%; Pred. No. 1.1e-41;
Matches 101; Conservative 52; Mismatches 112; Indels 10; Gaps 4;
QY 6 HSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDHSRRVPRTPWSSRISSQWLQL 65
DB 28 YSLTYITGLSKHVEDVPAPFQALGSLNDLQFFRYNSKDRKSQPMGLWRQVE-GMEDWKQD 86
QY 66 SQLKGDHMFVTVDFTIMENHNASKESHTLQVLGCMEQEDNSTEGYWKYGYDGODALE 125
DB 87 SQLQKAREDFMETLKDIVEYNDNSGSHVLOGRFGCEIENNRSSGAFWKYYDGDYIE 146
QY 126 FCPDITLWRAABPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGLVDOQVP 185
DB 147 ENKEIPAWVPDPAQITKQWEAEPPVYQARAYLEECFATLRKYLKYSKNILDRQDP 206
QY 186 PLVKVT-HHVTSSVTLRCALNYYPONTIMKWLKDKQPMDAKEFPK----DVLPGDGT 241
DB 207 PSVVVTSQAPGKCKKLCLAYDFYPGKIDVHWTAGEVQ-----EPRLRGDVLHNGNGT 261
QY 242 YQGWITLAVPPGEQRYTCQVEHPGLDPLIWIWE 276
DB 262 YQSWVVAVPPQDTAPYSCHVQHSSLAQPLVVPWE 296

RESULT 10
US-10-177-293-23
Sequence 23, Application US/10177293
Publication No. US200310124128A1
GENERAL INFORMATION:
APPLICANT: Lillie, James
APPLICANT: Glatt, Karen
APPLICANT: Zhao, Xumei
APPLICANT: Gannavarpu, Manjula
APPLICANT: Kamatkar, Shubhangi
APPLICANT: Mertens, Maureen
APPLICANT: Myer, Vic
APPLICANT: Wang, Youzhen
APPLICANT: Xu, Yongyao
APPLICANT: Hoersch, Sebastian
APPLICANT: Monahan, John
APPLICANT: Meyers, Rachel E.
APPLICANT: Bast Jr., Robert C.
APPLICANT: Hortobagyi, Gabriel N.
APPLICANT: Pusztai, Lajos
APPLICANT: Meric, Funda
APPLICANT: Sahin, Aysegul

APPLICANT: Mills, Gordon B.
TITLE OF INVENTION: COMPOSITIONS, KITS, AND METHODS FOR IDENTIFICATION, ASSESSMENT,
PREVENTION, AND THERAPY OF BREAST CANCER
FILE REFERENCE: MRI-038
CURRENT APPLICATION NUMBER: US/10/177,293
CURRENT FILING DATE: 2002-06-21
PRIOR APPLICATION NUMBER: US 60/299,887
PRIOR FILING DATE: 2001-06-21
PRIOR APPLICATION NUMBER: US 60/301,572
PRIOR FILING DATE: 2001-06-27
PRIOR APPLICATION NUMBER: US 60/306,501
PRIOR FILING DATE: 2001-07-18
PRIOR APPLICATION NUMBER: US 60/325,002
PRIOR FILING DATE: 2001-09-25
PRIOR APPLICATION NUMBER: US 60/362,585
PRIOR FILING DATE: 2002-03-05
PRIOR APPLICATION NUMBER: US 60/xxx,xxx
PRIOR FILING DATE: 2002-05-14
NUMBER OF SEQ ID NOS: 506
SOFTWARE: FastSeq for Windows Version 4.0
SEQ ID NO 23
LENGTH: 298
TYPE: PRT
ORGANISM: Homo sapiens
US-10-177-293-23

Query Match 32.1%; Score 486; DB 15; Length 298;
Best Local Similarity 36.7%; Pred. No. 1.1e-41;
Matches 101; Conservative 52; Mismatches 112; Indels 10; Gaps 4;
QY 6 HSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDHSRRVPRTPWSSRISSQWLQL 65
DB 28 YSLTYITGLSKHVEDVPAPFQALGSLNDLQFFRYNSKDRKSQPMGLWRQVE-GMEDWKQD 86
QY 66 SQLKGDHMFVTVDFTIMENHNASKESHTLQVLGCMEQEDNSTEGYWKYGYDGODALE 125
DB 87 SQLQKAREDFMETLKDIVEYNDNSGSHVLOGRFGCEIENNRSSGAFWKYYDGDYIE 146
QY 126 FCPDITLWRAABPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGLVDOQVP 185
DB 147 ENKEIPAWVPDPAQITKQWEAEPPVYQARAYLEECFATLRKYLKYSKNILDRQDP 206
QY 186 PLVKVT-HHVTSSVTLRCALNYYPONTIMKWLKDKQPMDAKEFPK----DVLPGDGT 241
DB 207 PSVVVTSQAPGKCKKLCLAYDFYPGKIDVHWTAGEVQ-----EPRLRGDVLHNGNGT 261
QY 242 YQGWITLAVPPGEQRYTCQVEHPGLDPLIWIWE 276
DB 262 YQSWVVAVPPQDTAPYSCHVQHSSLAQPLVVPWE 296

RESULT 11
US-10-015-535-32
Sequence 32, Application US/10015535
Publication No. US20030036506A1
GENERAL INFORMATION:
APPLICANT: Kranz, David M.
APPLICANT: Brophy, Susan
TITLE OF INVENTION: Mutated Class I Major Histocompatibility proteins and
Complexes
FILE REFERENCE: 100-00
CURRENT APPLICATION NUMBER: US/10/015,535
CURRENT FILING DATE: 2001-12-10
PRIOR APPLICATION NUMBER: 60/254,495
PRIOR FILING DATE: 2000-12-08
NUMBER OF SEQ ID NOS: 37
SOFTWARE: Patentin Ver. 2.0
SEQ ID NO 32
LENGTH: 542
TYPE: PRT
ORGANISM: Artificial Sequence
FEATURE:
OTHER INFORMATION: Description of Artificial Sequence: Synthetic


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Qy 182 QQVPLPVKVVTHV - TSSVTTLLRCRALNYYPNITMKMLKKQPMDAKEFFPKQVLPNGDG 24
Db 234 RTDSPKAVHTHSRSPEDKVTLLRCWALGFYPADIITLTQNGEEL - IQDMELVETRPA GDG 29
Qy 241 TYQGHITLAVPGBEQRVTCQVEHPGLDQPLVIWE 276
Db 293 TQKASVVVPLGKEQYVTCVTHQGLPEPLTRWE 328

Search completed: August 5, 2003, 13:21:56
Job time : 34 secs

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; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic
; ; OTHER INFORMATION: peptide
US-10-015-535-22

Query Match      30.9%; Score 468; DB 15; Length 540;
Best Local Similarity 39.5%; Pred. No. 1.7e-39;
Matches 109; Conservative 39; Mismatches 120; Indels 8; Gaps 7;

QY      6  HSLHYLFPMGLASEQDLGLSLFEALGYVDDQLFVFDH--ESRRVDPPTPTWSSRISSQMWL 63
      |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||  |||
Db      144 HSLRYFVTAVSRPGLGEPRIYMEVGYVDTEFVRFSDAENPRYEPFRARWMEQE-GPEYWE 202

QY      64  QLSQSLKGDHMFIVDPFWTMEHNHASK-SSHTLQVTLGCMEQEDNS-TGYWKYGYDQG 121
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db      203 RETQKAGNEQSPRVDLRTLLGYNQS KGSHTIQVISGCEVSGDGLLRGYQOYAYDGC 262

QY      122 DALFECPDITLDWAAEPPRAWPTKLEWHERHKIRARQNRAYLERDCPAQLQLLELGRGVLD 181
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db      263 DYALNEDLKTWTAADWAALITKHKEQAG-EAERLRAYLEGTCVEWLRRLKNGNATLL 321

QY      182 QQVPLPVKVTTHV-TGSVTTLRCLALNYYPQNTIMKWLKDQKPMDAKEBFPKVLNPGDG 240
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db      322 RTDSPKAHVTHSRSPEDKVTLCRWALGFYPADITLTQLNGEEL-IQDMELVETRPAGDG 380

QY      241 TYQGWITLAVPPGEEQRYTCQVHPGLDQPLIWIWE 276
      : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : : :
Db      381 TFOKASVVPVLGKEQYITCVHQGLPEPLTLRW 416

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RESULT 15
US-10-015-535-28
; Sequence 28, Application US/10015535
; Publication No. US20030036506A1
; GENERAL INFORMATION:
; APPLICANT: Kranz, David M.
; APPLICANT: Brophy, Susan
; TITLE OF INVENTION: Mutated Class I Major Histocompatibility proteins and
; TITLE OF INVENTION: Complexes
; FILE REFERENCE: 100-00
; CURRENT APPLICATION NUMBER: US/10/015,535
; CURRENT FILING DATE: 2001-12-10
; PRIOR APPLICATION NUMBER: 60/254,495
; PRIOR FILING DATE: 2000-12-08
; NUMBER OF SEQ ID NOS: 37
; SOFTWARE: PatentIn Ver. 2.0
; SEQ ID NO 28
; LENGTH: 541
; TYPE: PRT
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Description of Artificial Sequence: Synthetic
; OTHER INFORMATION: peptide
US-10-015-535-28

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Query Match	30.9%	Score 468	DB 15	Length 541
Best Local Similarity	39.5%	Pred. No. 1.7e-39		
Matches 109; Conservative	39	Mismatches 120	Indels 8	Gaps 7

QY	6	HSLSHLYFMGASEQDILGLSLFEALGYVDQQLFVFDH--ESRRVEPTPTPVSSRISSQMWL	63
DB	56	HSLSRYFTVAVSRCLEGEPRYMEYGVDDTEFVFDSDAENPRVEPRARMWEQE-GPEYWE	114
QY	64	QLSLSKLGMDHMTFVDFWTIMENHNASK--ESHTLQVILGCEMOEDNS--TEGYWKYGYDQG	121
DB	115	RETQAKAGNEQSFRVLDRTLGLYNGSKGSSHTIQVISGCEVSDGELLRGYQQYAYDGC	174
QY	122	DALEFCDTLDWAAEPRAPWPTKLEWHRKIRARONRAYLERDCPAQLQLLELGRGVLD	181
DB	175	DYVALNEDLKTWTAADMAALITKHWEQAG-EAERLRAYLEGTCEVWRLRYLKNGNATLL	233

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:07:04 ; Search time 14.5 Seconds
(without alignments)
805.365 Million cell updates/sec

Title: US-10-092-404-3
Perfect score: 1514
Sequence: 1 RLRSLSHLFLWGASEQDL.....RYTCQVEHPLDQPLIVWE 276

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 328717 seqs, 4231058 residues

Total number of hits satisfying chosen parameters: 328717

Minimum DB seq length: 0
Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 45 summaries

Database : Issued Patents AA:*
1: /cgn2_6/ptodata/1/iaa/5A COMB.pep.*
2: /cgn2_6/ptodata/1/iaa/5B COMB.pep.*
3: /cgn2_6/ptodata/1/iaa/6A COMB.pep.*
4: /cgn2_6/ptodata/1/iaa/6B COMB.pep.*
5: /cgn2_6/ptodata/1/iaa/PCTUS COMB.pep.*
6: /cgn2_6/ptodata/1/iaa/backfile1.pep.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1514	100.0	276	4	US-09-094-964-3
2	1502	99.2	276	4	US-09-094-964-1
3	1502	99.2	348	3	US-08-652-265-2
4	1502	99.2	348	3	US-08-834-497A-2
5	1502	99.2	348	3	US-09-503-444A-2
6	1502	99.2	348	4	US-09-277-457-2
7	1502	99.2	348	4	US-09-679-729-2
8	1493	98.6	276	4	US-09-094-964-2
9	1493	98.6	348	3	US-08-652-265-6
10	1493	98.6	348	3	US-08-834-497A-6
11	1491	98.6	348	3	US-09-503-444A-6
12	1491	98.5	348	3	US-08-652-265-4
13	1491	98.5	348	3	US-08-834-497A-4
14	1491	98.5	348	3	US-09-503-444A-4
15	1482	97.9	348	3	US-08-652-265-8
16	1482	97.9	348	3	US-08-834-497A-8
17	1482	97.9	348	3	US-09-503-444A-8
18	517	34.1	361	3	US-08-652-265-22
19	517	34.1	361	3	US-08-834-497A-22
20	517	34.1	361	3	US-09-503-444A-22
21	511	33.8	364	4	US-08-914-372C-11
22	508	33.6	365	3	US-08-652-265-23
23	508	33.6	365	3	US-08-834-497A-23
24	508	33.6	365	3	US-09-503-444A-23
25	500	33.0	274	2	US-08-484-905-107
26	500	33.0	274	3	US-08-481-985B-107
27	500	33.0	274	3	US-08-370-476-107

28	500	33.0	341	3	US-08-890-719-38	Sequence 38, Appl
29	499	33.0	365	2	US-08-484-905-97	Sequence 97, Appl
30	499	33.0	365	3	US-08-481-985B-97	Sequence 97, Appl
31	499	33.0	365	3	US-08-370-476-97	Sequence 97, Appl
32	498	32.9	274	2	US-08-484-905-108	Sequence 108, App
33	498	32.9	274	3	US-08-481-985B-108	Sequence 108, App
34	498	32.9	274	3	US-08-370-476-108	Sequence 108, App
35	498	32.9	365	3	US-08-484-905-100	Sequence 100, App
36	498	32.9	365	3	US-08-481-985B-100	Sequence 100, App
37	498	32.9	365	3	US-08-370-476-100	Sequence 100, App
38	497	32.8	274	1	US-08-222-851-1	Sequence 1, Appli
39	497	32.8	363	4	US-08-914-372C-37	Sequence 37, Appl
40	497	32.8	365	2	US-08-484-905-99	Sequence 99, Appl
41	497	32.8	365	3	US-08-481-985B-99	Sequence 99, Appl
42	497	32.8	365	3	US-08-370-476-99	Sequence 99, Appl
43	496	32.8	274	2	US-08-484-905-106	Sequence 106, App
44	496	32.8	274	3	US-08-481-985B-106	Sequence 106, App
45	496	32.8	274	3	US-08-370-476-106	Sequence 106, App

ALIGNMENTS

RESULT 1
US-09-094-964-3
; Sequence 3, Application US/09094964
; Patent No. 6391852
; GENERAL INFORMATION:
; APPLICANT: Feder, John N.
; APPLICANT: Bjorkman, Pamela J.
; APPLICANT: Schatzman, Randall C.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
; TITLE OF INVENTION: DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
; TITLE OF INVENTION: AND IRON DEFICIENCY DISEASES
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds, LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10036-2811
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; OPERATING SYSTEM: Windows
; SOFTWARE: FastSeq for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/094,964
; FILING DATE: June 12, 1998
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/876,010
; FILING DATE: June 13, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0074-999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-493-4935
; TELEFAX: 650-493-5556
; TELEX: 66141 PENNIE
; INFORMATION FOR SEQ ID NO: 3:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 276 amino acids
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
; US-09-094-964-3
Query Match 100.0%; Score 1514; DB 4; Length 276;
Best Local Similarity 100.0%; Pred. No. 6.2e-144;

	Matches	276;	Conservative	0;	Mismatches	0;	Indels	0;	Gaps	0;
Qy	1	RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFYDHSRRRVEPRTPWSSRISSQ	60							
Db	1	RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFYDHSRRRVEPRTPWSSRISSQ	60							
Qy	61	MWLQSLSKGDWHFTVDFWTIMENHNASKESHTLQVLGCMEQDNSTEGYWKYGVDG	120							
Db	61	MWLQSLSKGDWHFTVDFWTIMENHNASKESHTLQVLGCMEQDNSTEGYWKYGVDG	120							
Qy	121	QDALEFCPDTLDWRAAEPRAWPTKLWEHRHKIRARQNAYLERDCPAOLOLLBLGRGVL	180							
Db	121	QDALEFCPDTLDWRAAEPRAWPTKLWEHRHKIRARQNAYLERDCPAOLOLLBLGRGVL	180							
Qy	181	DQQVPPLVKVTHHVTSSVTTILRCRALNYYPONITMKWLKDQPMDAKEFEPKDVLPNGDG	240							
Db	181	DQQVPPLVKVTHHVTSSVTTILRCRALNYYPONITMKWLKDQPMDAKEFEPKDVLPNGDG	240							
Qy	241	TYQGWI TLAVPPGEOR YTCOVERH PGLDQPLIV IWE	276							
Db	241	TYQGWI TLAVPPGEOR YTCOVERH PGLDQPLIV IWE	276							

RESULT 2
 US-09-094-964-1
 Sequence 1, Application US/09094964
 Patent No. 6391852
 GENERAL INFORMATION:
 APPLICANT: Feder, John N.
 APPLICANT: Bjorkman, Pamela J.
 APPLICANT: Schatzman, Randall C.
 TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
 TITLE OF INVENTION: DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
 TITLE OF INVENTION: DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
 NUMBER OF SEQUENCES: 5
 CORRESPONDENCE ADDRESS:
 ADDRESSEE: Pennie & Edmonds, LLP
 STREET: 1155 Avenue of the Americas
 CITY: New York
 STATE: NY
 COUNTRY: USA
 ZIP: 10036-2811
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Diskette
 COMPUTER: IBM Compatible
 OPERATING SYSTEM: Windows
 SOFTWARE: FastSEQ for Windows Version 2.0b
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/09/094,964
 FILING DATE: June 12, 1998
 CLASSIFICATION:
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: 08/876,010
 FILING DATE: June 13, 1997
 ATTORNEY/AGENT INFORMATION:
 NAME: Poissant, Brian M
 REGISTRATION NUMBER: 28,462
 REFERENCE/DOCKET NUMBER: 8907-0074-999
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: 650-493-4935
 TELEFAX: 650-493-5556
 TELEX: 66141 PENNIE
 INFORMATION FOR SEQ ID NO: 1:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 276 amino acids
 TYPE: amino acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: peptide
 US-09-094-964-1

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Matches 274; Conservative 0; Mismatches 2; Indels

Qy 1 RLLRSHSLHYLFMGASEQDGLGLSLFEALGYVDDQLFVYDHESRRVRRPRT
Db 1 RLLRSHSLHYLFMGASEQDGLGLSLFEALGYVDDQLFVYDHESRRVRRPRT
Qy 61 MWLQLSQSLKGDWHMFTVDFWTIMENHNASKESHTLQVILGCENQEDNST
Db 61 MWLQLSQSLKGDWHMFTVDFWTIMENHNASKESHTLQVILGCENQEDNST
Qy 121 QDALEFCPDTLDWRAAEPRAPWPKLEWERHKIRARQNRAVLERDCPAQLQ
Db 121 QDHLEFCPDTLDWRAAEPRAPWPKLEWERHKIRARQNRAVLERDCPAQLQ
Qy 181 DQQVPLVKVTHVTSVVTILRCALNYYQNTITMKWLKDKQPMDAKEFE
Db 181 DQQVPLVKVTHVTSVVTILRCALNYYQNTITMKWLKDKQPMDAKEFE
Qy 241 TYQGWTITLAVPPGEQRYTCQVHPGLDQPLIWIWE 276
Db 241 TYQGWTITLAVPPGEQRYTCQVHPGLDQPLIWIWE 276

RESULT 3
US-08-652-265-2
; Sequence 2, Application US/08652265
; Patent No. 6025130
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchinashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: Hereditary Hemochromatosis Gene
; NUMBER OF SEQUENCES: 44
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew LLP
; STREET: Two Embarcadero Center, Eighth Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/652,265
; FILING DATE: 23-MAY-1996
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M.
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 17957-000500
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 576-0200
; TELEFAX: (415) 576-0300
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-652-265-2

Query Match 99.2%; Score 1502; DB 3; Length 348
Best Local Similarity 99.3%; Pred. No. 1.4e-142;
Matches 274; Conservative 0; Mismatches 2; Indels

Qy 1 RLLRSHSLHYLFMGASEQDGLGLSLFEALGYVDDQLFVYDHESRRVRRPRT

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Db 23 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDQLFVFDHESRRVPRTPWVSSRISSQ 82
QY 61 MWLQLSQSLKGDHMTVDFTWIMENHNASKESHTLQVLGCEMOEDNSTEGYWKYGYDG 120
Db 83 MWLQLSQSLKGDHMTVDFTWIMENHNASKESHTLQVLGCEMOEDNSTEGYWKYGYDG 142
QY 121 QDALFECPTDLDWRAAEPRAMPKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
Db 143 QDHLEFCPTDLDWRAAEPRAMPKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DOQVPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
Db 203 DOQVPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHFGLDQPLIVWE 276
Db 263 TYQGWITLAVPPGEEQRYTCQVEHFGLDQPLIVWE 298

RESULT 4

US-08-834-497A-2
; Sequence 2, Application US/08834497A
; Patent No. 6140305

; GENERAL INFORMATION:

; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: HEREDITARY HEMOCHROMATOSIS GENE PRODUCTS
; NUMBER OF SEQUENCES: 76
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036-2811

; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: FASTSEQ for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/834,497A
; FILING DATE: 04-APR-1997

; CLASSIFICATION: 514

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: US 08/652,265

; FILING DATE: 23-MAY-1996

; CLASSIFICATION: 514

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: US 08/632,673

; FILING DATE: 16-APR-1996

; CLASSIFICATION: 514

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: US 08/630,912

; FILING DATE: 04-APR-1996

; CLASSIFICATION: 514

; ATTORNEY/AGENT INFORMATION:

; NAME: Poissant, Brian M.

; REGISTRATION NUMBER: 28,462

; REFERENCE/DOCKET NUMBER: 8907-0056-999

; TELECOMMUNICATION INFORMATION:

; TELEPHONE: 650-493-4935

; TELEFAX: 650-493-5556

; TELEX: 66141 PENNIE

; INFORMATION FOR SEQ ID NO: 2:

; SEQUENCE CHARACTERISTICS:

; LENGTH: 348 amino acids

; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-834-497A-2

Query Match 99.2%; Score 1502; DB 3; Length 348;

Best Local Similarity 99.3%; Pred. No. 1.4e-142;

Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDQLFVFDHESRRVPRTPWVSSRISSQ 60

Db 23 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDQLFVFDHESRRVPRTPWVSSRISSQ 82

QY 61 MWLQLSQSLKGDHMTVDFTWIMENHNASKESHTLQVLGCEMOEDNSTEGYWKYGYDG 120

Db 83 MWLQLSQSLKGDHMTVDFTWIMENHNASKESHTLQVLGCEMOEDNSTEGYWKYGYDG 142

QY 121 QDALFECPTDLDWRAAEPRAMPKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180

Db 143 QDHLEFCPTDLDWRAAEPRAMPKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202

QY 181 DOQVPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240

Db 203 DOQVPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 262

QY 241 TYQGWITLAVPPGEEQRYTCQVEHFGLDQPLIVWE 276

Db 263 TYQGWITLAVPPGEEQRYTCQVEHFGLDQPLIVWE 298

RESULT 5

US-09-503-444A-2

; Sequence 2, Application US/09503444A

; Patent No. 6228594

; GENERAL INFORMATION:

; APPLICANT: Thomas, Winston J.

; APPLICANT: Drayna, Dennis T.

; APPLICANT: Feder, John N.

; APPLICANT: Gnirke, Andreas

; APPLICANT: Ruddy, David

; APPLICANT: Tsuchihashi, Zenta

; APPLICANT: Wolff, Roger K.

; TITLE OF INVENTION: Hereditary Hemochromatosis Gene

; NUMBER OF SEQUENCES: 44

; CORRESPONDENCE ADDRESS:

; ADDRESSEE: Pennie & Edmonds LLP

; STREET: 1155 Avenue of the Americas

; CITY: New York

; STATE: New York

; COUNTRY: USA

; ZIP: 10036

; COMPUTER READABLE FORM:

; MEDIUM TYPE: Floppy disk

; COMPUTER: IBM PC compatible

; OPERATING SYSTEM: Windows 95

; SOFTWARE: WordPerfect Version 8

; CURRENT APPLICATION DATA:

; APPLICATION NUMBER: US/09/503,444A

; FILING DATE: 14-Feb-2000

; CLASSIFICATION:

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: 08/652,265

; FILING DATE: 23-May-1996

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: 08/632,673

; FILING DATE: 16-Apr-1996

; PRIOR APPLICATION DATA:

; APPLICATION NUMBER: 08/630,912

; FILING DATE: 04-Apr-1996

; ATTORNEY/AGENT INFORMATION:

; NAME: Poissant, Brian M.

; REGISTRATION NUMBER: 28,462

; REFERENCE/DOCKET NUMBER: 8907-0088-999

TELECOMMUNICATION INFORMATION:

TELEPHONE: 212-790-9090

TELEFAX: 212-869-9741

TELEX: 66141

INFORMATION FOR SEQ ID NO: 2:

SEQUENCE CHARACTERISTICS:

LENGTH: 348 amino acids

TYPE: amino acid

TOPOLOGY: linear

MOLECULE TYPE: protein

US-09-503-444A-2

Query Match 99.2%; Score 1502; DB 3; Length 348;

Best Local Similarity 99.3%; Pred. No. 1.4e-142;

Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISQ 60
DB 23 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISQ 82
QY 61 MWLQLSQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPDTLDWRAAEPRAWPTKLEWERHKIRARQNRAYLERDCCPAQLQQLLELGRGVL 180
DB 143 QDHLEFCPDTLDWRAAEPRAWPTKLEWERHKIRARQNRAYLERDCCPAQLQQLLELGRGVL 202
QY 181 DQOVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240
DB 203 DQOVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 262
QY 241 TYQGWITLAVPGEORQYTCQVEHGLDQPLIVWE 276
DB 263 TYQGWITLAVPGEORQYTCQVEHGLDQPLIVWE 298

RESULT 6

US-09-277-457-2

Sequence 2, Application US/09277457

Patent No. 6355425

GENERAL INFORMATION:

APPLICANT: Rothenberg, Barry E.

APPLICANT: Sawada-Hirai, Ritsuko

APPLICANT: Barton, James C.

TITLE OF INVENTION: MUTATIONS ASSOCIATED WITH IRON DISORDERS

FILE REFERENCE: 10653/002001

CURRENT APPLICATION NUMBER: US/09/277,457

CURRENT FILING DATE: 1999-03-26

NUMBER OF SEQ ID NOS: 30

SOFTWARE: FastSeq for Windows Version 4.0

SEQ ID NO 2

LENGTH: 348

TYPE: PRT

ORGANISM: Homo Sapiens

US-09-277-457-2

Query Match

Best Local Similarity 99.2%; Score 1502; DB 4; Length 348;

Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISQ 60
DB 23 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISQ 82
QY 61 MWLQLSQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPDTLDWRAAEPRAWPTKLEWERHKIRARQNRAYLERDCCPAQLQQLLELGRGVL 180
DB 143 QDHLEFCPDTLDWRAAEPRAWPTKLEWERHKIRARQNRAYLERDCCPAQLQQLLELGRGVL 202

QY 181 DQOVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240
DB 203 DQOVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 262
QY 241 TYQGWITLAVPGEORQYTCQVEHGLDQPLIVWE 276
DB 263 TYQGWITLAVPGEORQYTCQVEHGLDQPLIVWE 298

RESULT 7

US-09-679-729-2

Sequence 2, Application US/09679729

Patent No. 6509442

GENERAL INFORMATION:

APPLICANT: Rothenberg, Barry E.

APPLICANT: Sawada-Hirai, Ritsuko

APPLICANT: Barton, James C.

TITLE OF INVENTION: MUTATIONS ASSOCIATED WITH IRON DISORDERS

FILE REFERENCE: 24065-004 DIV

CURRENT APPLICATION NUMBER: US/09/679,729

CURRENT FILING DATE: 2000-10-04

PRIOR APPLICATION NUMBER: 09/277,457

PRIOR FILING DATE: 1999-03-26

NUMBER OF SEQ ID NOS: 30

SOFTWARE: FastSeq for Windows Version 4.0

SEQ ID NO 2

LENGTH: 348

TYPE: PRT

ORGANISM: Homo Sapiens

US-09-679-729-2

Query Match

Best Local Similarity 99.2%; Score 1502; DB 4; Length 348;

Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISQ 60
DB 23 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISQ 82
QY 61 MWLQLSQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGWDHMTVDFTWMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPDTLDWRAAEPRAWPTKLEWERHKIRARQNRAYLERDCCPAQLQQLLELGRGVL 180
DB 143 QDHLEFCPDTLDWRAAEPRAWPTKLEWERHKIRARQNRAYLERDCCPAQLQQLLELGRGVL 202
QY 181 DQOVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 240
DB 203 DQOVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGD 262
QY 241 TYQGWITLAVPGEORQYTCQVEHGLDQPLIVWE 276
DB 263 TYQGWITLAVPGEORQYTCQVEHGLDQPLIVWE 298

RESULT 8

US-09-094-964-2

Sequence 2, Application US/09094964

Patent No. 6391852

GENERAL INFORMATION:

APPLICANT: Feder, John N.

APPLICANT: Bjorkman, Pamela J.

APPLICANT: Schatzman, Randall C.

TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR

TITLE OF INVENTION: DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES

NUMBER OF SEQUENCES: 5

CORRESPONDENCE ADDRESS:

ADDRESSEE: Pennie & Edmonds, LLP

STREET: 1155 Avenue of the Americas

CITY: New York

STATE: NY

COUNTRY: USA
ZIP: 10036-2811
COMPUTER READABLE FORM:
MEDIUM TYPE: Diskette
COMPUTER: IBM Compatible
OPERATING SYSTEM: Windows
SOFTWARE: FASTSO for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/094,964
FILING DATE: June 12, 1998
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/876,010
FILING DATE: June 13, 1997
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M
REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0074-999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 276 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-094-964-2

Query Match 98.6%; Score 1493; DB 4; Length 276;
Best Local Similarity 98.9%; Pred. No. 7.9e-142;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLRSHSLHYLFMGASEODLGLSFEALGYVDDQLFVFDHESRRVPRTPWSSRISQ 60
DB 1 RLRSHSLHYLFMGASEODLGLSFEALGYVDDQLFVFDHESRRVPRTPWSSRISQ 60
QY 61 MWLQSQSLKGDHMTVDFTWIMENHNASKESHTLQVILGCEQEDNSTEGYWKYGYDG 120
DB 61 MWLQSQSLKGDHMTVDFTWIMENHNASKESHTLQVILGCEQEDNSTEGYWKYGYDG 120
QY 121 QDALEFCPTLDWRAAPRAWPTKLEWERHKIRARQRAYLERDCAQLQQLLELGRGVL 180
DB 121 QDHLFCPTLDWRAAPRAWPTKLEWERHKIRARQRAYLERDCAQLQQLLELGRGVL 180
QY 181 DQOVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
DB 181 DQOVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
QY 241 TYQGWITLAVPGEORVTCQVEHPGLDQPLIVWE 276
DB 241 TYQGWITLAVPGEORVTCQVEHPGLDQPLIVWE 276

RESULT 9
US-08-652-265-6
Sequence 6, Application US/08652265
Patent No. 6025130
GENERAL INFORMATION:
APPLICANT: Thomas, Winston J.
APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: Hereditary Hemochromatosis Gene
NUMBER OF SEQUENCES: 44
CORRESPONDENCE ADDRESS:
ADDRESSEE: Townsend and Townsend and Crew LLP
STREET: Two Embarcadero Center, Eighth Floor

CITY: San Francisco
STATE: California
COUNTRY: USA
ZIP: 94111-3834
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/652,265
FILING DATE: 23-MAY-1996
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Smith, William M.
REGISTRATION NUMBER: 30,223
REFERENCE/DOCKET NUMBER: 17957-000500
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 576-0200
TELEFAX: (415) 576-0300
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-652-265-6
Query Match 98.6%; Score 1493; DB 3; Length 348;
Best Local Similarity 98.9%; Pred. No. 1.1e-141;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLRSHSLHYLFMGASEODLGLSFEALGYVDDQLFVFDHESRRVPRTPWSSRISQ 60
DB 23 RLRSHSLHYLFMGASEODLGLSFEALGYVDDQLFVFDHESRRVPRTPWSSRISQ 82
QY 61 MWLQSQSLKGDHMTVDFTWIMENHNASKESHTLQVILGCEQEDNSTEGYWKYGYDG 120
DB 83 MWLQSQSLKGDHMTVDFTWIMENHNASKESHTLQVILGCEQEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPTLDWRAAPRAWPTKLEWERHKIRARQRAYLERDCAQLQQLLELGRGVL 180
DB 143 QDHLFCPTLDWRAAPRAWPTKLEWERHKIRARQRAYLERDCAQLQQLLELGRGVL 202
QY 181 DQOVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
DB 203 DQOVPLVKVTHVTSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 262
QY 241 TYQGWITLAVPGEORVTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPGEORVTCQVEHPGLDQPLIVWE 298
RESULT 10
US-08-834-497A-6
Sequence 6, Application US/08834497A
Patent No. 6140305
GENERAL INFORMATION:
APPLICANT: Thomas, Winston J.
APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: HEREDITARY HEMOCHROMATOSIS GENE PRODUCTS
NUMBER OF SEQUENCES: 76
CORRESPONDENCE ADDRESS:
ADDRESSEE: pennie & Edmonds LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA

ZIP: 10036-2811
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: FASTSEQ for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/834,497A
FILING DATE: 04-APR-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/652,265
FILING DATE: 23-MAY-1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/632,673
FILING DATE: 16-APR-1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/630,912
FILING DATE: 04-APR-1996
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M.
REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0056-999
TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-834-497A-6

Query Match 98.6%; Score 1493; DB 3; Length 348;
Best Local Similarity 98.9%; Pred. No. 1.1e-141;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLRLSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWVSSRISSQ 60
DB 23 RLRLSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWVSSRISSQ 82
QY 61 MWLQSLQSLKGDHMTFTVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGDG 120
DB 83 MWLQSLQSLKGDHMTFTVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGDG 142
QY 121 QDALRCPDPTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLEFCPTDLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQVPPPLVKVTHVTSVTTLCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGNGD 240
DB 203 DQVPPPLVKVTHVTSVTTLCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGNGD 262
QY 241 TYQGWITLAVPGEQRYTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPGEQRYTCQVEHPGLDQPLIVWE 298

RESULT 11
US-09-503-444A-6
Sequence 6, Application US/09503444A
Patent No. 6228594
GENERAL INFORMATION:
APPLICANT: Thomas, Winston J.
APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David

APPLICANT: Teuchiashashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: Hereditary Hemochromatosis Gene
NUMBER OF SEQUENCES: 44
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: WordPerfect version 8
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/503,444A
FILING DATE: 14-Feb-2000
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/652,265
FILING DATE: 23-May-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/632,673
FILING DATE: 16-Apr-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/630,912
FILING DATE: 04-Apr-1996
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M.
REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0088-999
TELEPHONE: 212-790-9090
TELEFAX: 212-869-9741
TELEX: 66141
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-503-444A-6
Query Match 98.6%; Score 1493; DB 3; Length 348;
Best Local Similarity 98.9%; Pred. No. 1.1e-141;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLRLSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWVSSRISSQ 60
DB 23 RLRLSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWVSSRISSQ 82
QY 61 MWLQSLQSLKGDHMTFTVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGDG 120
DB 83 MWLQSLQSLKGDHMTFTVDFWTIMENHNASKESHTLQVILGCEMQEDNSTEGYWKYGDG 142
QY 121 QDALRCPDPTLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLEFCPTDLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQVPPPLVKVTHVTSVTTLCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGNGD 240
DB 203 DQVPPPLVKVTHVTSVTTLCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGNGD 262
QY 241 TYQGWITLAVPGEQRYTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPGEQRYTCQVEHPGLDQPLIVWE 298

RESULT 12
US-08-652-265-4
Sequence 4, Application US/08652265

Patent No. 6025130
GENERAL INFORMATION:
APPLICANT: Thomas, Winston J.
APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: Hereditary Hemochromatosis Gene
NUMBER OF SEQUENCES: 44
CORRESPONDENCE ADDRESS:
ADDRESSEE: Townsend and Townsend and Crew LLP
STREET: Two Embarcadero Center, Eighth Floor
CITY: San Francisco
STATE: California
COUNTRY: USA
ZIP: 94111-3834
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/652,265
FILING DATE: 23-MAY-1996
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Smith, William M.
REGISTRATION NUMBER: 30,223
REFERENCE/DOCKET NUMBER: 17957-000500
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 576-0200
TELEFAX: (415) 576-0300
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-652-265-4

Query Match 98.5%; Score 1491; DB 3; Length 348;
Best Local Similarity 98.9%; Pred. No. 1.7e-141;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISQ 60
DB 23 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISQ 82
QY 61 MWLQSLQSLKGWDMFTVDFTWMENHNHNSKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGWDMFTVDFTWMENHNHNSKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPDTLDWRAAEPRAMPPTKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGYL 180
DB 143 QDHLFCPDTLDWRAAEPRAMPPTKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGYL 202
QY 181 DOQVPLVKVTHHTVSSVTLTLCRALNYPONITMKWLKDKQPMDAKEPEPKDVLPGD 240
DB 203 DOQVPLVKVTHHTVSSVTLTLCRALNYPONITMKWLKDKQPMDAKEPEPKDVLPGD 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 298

RESULT 13

US-08-834-497A-4
Sequence 4, Application US/08834497A
Patent No. 6140305
GENERAL INFORMATION:
APPLICANT: Thomas, Winston J.

APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: HEREDITARY HEMOCHROMATOSIS GENE PRODUCTS
NUMBER OF SEQUENCES: 76
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036-2811
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: FastSeq for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/834,497A
FILING DATE: 04-APR-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/652,265
FILING DATE: 23-MAY-1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/632,673
FILING DATE: 16-APR-1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/630,912
FILING DATE: 04-APR-1996
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M.
REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0056-999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 4:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-834-497A-4
Query Match 98.5%; Score 1491; DB 3; Length 348;
Best Local Similarity 98.9%; Pred. No. 1.7e-141;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISQ 60
DB 23 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDDQLFVFDHESRRRVEPRTPWSSRISQ 82
QY 61 MWLQSLQSLKGWDMFTVDFTWMENHNHNSKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGWDMFTVDFTWMENHNHNSKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPDTLDWRAAEPRAMPPTKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGYL 180
DB 143 QDHLFCPDTLDWRAAEPRAMPPTKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGYL 202
QY 181 DOQVPLVKVTHHTVSSVTLTLCRALNYPONITMKWLKDKQPMDAKEPEPKDVLPGD 240
DB 203 DOQVPLVKVTHHTVSSVTLTLCRALNYPONITMKWLKDKQPMDAKEPEPKDVLPGD 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276

Db 263 TYQGWITLAVPGEQRYTQVEHPGLDQPLIWIWE 298
|||||
RESULT 14
US-09-503-444A-4
; Sequence 4, Application US/09503444A
; Patent No. 6228594
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: Hereditary Hemochromatosis Gene
; NUMBER OF SEQUENCES: 44
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: WordPerfect Version 8
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/503,444A
; FILING DATE: 14-Feb-2000
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/652,265
; FILING DATE: 23-May-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/632,673
; FILING DATE: 16-Apr-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/630,912
; FILING DATE: 04-Apr-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M.
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0088-999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 212-790-9090
; TELEFAX: 212-869-9741
; TELEX: 66141
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-09-503-444A-4
Query Match 98.5%; Score 1491; DB 3; Length 348;
Best Local Similarity 98.9%; Pred. No. 1.7e-141;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISQ 60
Db 23 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISQ 82
QY 61 MWLQLSQSLKGWDHMTFTVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDG 120
Db 83 MWLQLSQSLKGWDHMTFTVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPTDLWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGYL 180
Db 143 QHLEFCPTDLWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGYL 202
QY 181 DOQVPLVKVTHVTSSVTTLCRALNYYPQNTWKWLKDKQPMDAKEPEPKDVLNPGDG 240
|||||

Db 143 QDHLFCPTDLWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGYL 202
QY 181 DOQVPLVKVTHVTSSVTTLCRALNYYPQNTWKWLKDKQPMDAKEPEPKDVLNPGDG 240
Db 203 DOQVPLVKVTHVTSSVTTLCRALNYYPQNTWKWLKDKQPMDAKEPEPKDVLNPGDG 262
QY 241 TYQGWITLAVPGEQRYTQVEHPGLDQPLIWIWE 276
Db 263 TYQGWITLAVPGEQRYTQVEHPGLDQPLIWIWE 298
|||||

RESULT 15
US-08-652-265-8
; Sequence 8, Application US/08652265
; Patent No. 6025130
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: Hereditary Hemochromatosis Gene
; NUMBER OF SEQUENCES: 44
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew LLP
; STREET: Two Embarcadero Center, Eighth Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/652,265
; FILING DATE: 23-MAY-1996
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M.
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 17957-000500
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 576-0200
; TELEFAX: (415) 576-0300
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-652-265-8
Query Match 97.9%; Score 1482; DB 3; Length 348;
Best Local Similarity 98.6%; Pred. No. 1.4e-140;
Matches 272; Conservative 0; Mismatches 4; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISQ 60
Db 23 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVFDHESRRVPRTPWSSRISQ 82
QY 61 MWLQLSQSLKGWDHMTFTVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDG 120
Db 83 MWLQLSQSLKGWDHMTFTVDFWTIMENHNASKESHTLQVILGCEMOEDNSTEGYWKYGYDG 142
QY 121 QDALEFCPTDLWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGYL 180
Db 143 QHLEFCPTDLWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGYL 202
QY 181 DOQVPLVKVTHVTSSVTTLCRALNYYPQNTWKWLKDKQPMDAKEPEPKDVLNPGDG 240
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||||| 203 DQVPPLVKVTHVTSSVTLRCRALNYFPQNTMKWLKDKQPMDAKEFEFKDVLPNGDG 262
Qy      241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
||||| 263 TYQGWITLAVPPGEEQRYTYQVEHPGLDQPLIWIWE 298
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Search completed: August 5, 2003, 13:11:12
Job time : 15.5 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:05:29 ; Search time 38 Seconds
(without alignments)

1152.856 Million cell updates/sec

Title: US-10-092-404-2

Perfect score: 1520

Sequence: 1 RLRLSHLHLYFLMGASEQDL.....RYTCQVHPGLDPLIVWE 276

Scoring table: BLOSUM62

Gapop 10.0 , Gapext 0.5

Searched: 1107863 seqs, 158726573 residues

Total number of hits satisfying chosen parameters: 1107863

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	1520	100.0	276	20 AAW94296	HFE mutant (H63D-H
2	1520	100.0	276	24 ABG72686	Human haemochromat
3	1520	100.0	348	22 AAB36871	Human hereditary h
4	1513	99.5	276	20 AAW94295	Wild-type HFE poly
5	1513	99.5	276	24 ABG72685	Human haemochromat
6	1513	99.5	348	18 AAW36499	Hereditary haemoch
7	1513	99.5	348	21 AAB19149	A human histocoma
8	1513	99.5	348	22 AAB36869	Human hereditary h
9	1509	99.3	348	22 AAB36872	Human hereditary h

10	1508	99.2	438	23 AAU00035	Beta 2 microglobul
11	1502	98.8	348	22 AAB36870	Human hereditary h
12	1493	98.2	276	20 AAW94297	HFE mutant (H111A/
13	1493	98.2	276	24 ABG72687	Human haemochromat
14	523	34.4	361	22 AAB36873	Rabbit leukocyte a.
15	514	33.8	92	24 ABF68379	Human colon specif
16	514	33.8	365	22 AAB36874	MHC class I protei
17	506	33.3	274	21 AAY68275	Human leukocyte an
18	506	33.3	274	21 AAY52929	HLA-A2/A28 family
19	506	33.3	274	22 AAB58690	HLA-A2/A28 protein
20	506	33.3	280	22 ABU10225	Human leukocyte an
21	506	33.3	280	24 ABU08672	Human histocompat
22	506	33.3	415	22 ABU10224	Human partial beta
23	506	33.3	415	24 ABU08671	Human single chain
24	505	33.2	365	21 AAY68265	Human leukocyte an
25	505	33.2	365	21 AAY52919	HLA-A2/A28 family
26	505	33.2	365	22 AAB58680	Human leukocyte an
27	505	33.2	368	22 AAM24017	Human EST encoded
28	504	33.2	274	21 AAY68276	Human leukocyte an
29	504	33.2	274	21 AAY52930	HLA-A2/A28 family
30	504	33.2	274	22 AAB58691	HLA-A2/A28 protein
31	504	33.2	365	21 AAY68268	Human leukocyte an
32	504	33.2	365	21 AAY52922	HLA-A2/A28 family
33	504	33.2	365	22 AAB58683	HLA-A2/A28 protein
34	503	33.1	274	9 AAP80911	Consensus sequence
35	503	33.1	365	21 AAY68267	Human leukocyte an
36	503	33.1	365	21 AAY52921	HLA-A2/A28 family
37	503	33.1	365	22 AAB58682	HLA-A2/A28 protein
38	502	33.0	274	21 AAY68274	Human leukocyte an
39	502	33.0	274	21 AAY52928	HLA-A2/A28 family
40	502	33.0	274	22 AAB58689	HLA-A2/A28 protein
41	502	33.0	365	21 AAY68266	Human leukocyte an
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43	502	33.0	365	22 AAB58681	HLA-A2/A28 protein
44	501	33.0	412	19 AAW68385	Chimeric HLA-A2.1/
45	500	32.9	274	21 AAY68273	Human leukocyte an

ALIGNMENTS

RESULT 1

AAW94296

ID AAW94296 standard; peptide; 276 AA.

XX AC

XX AAW94296;

XX DT

DT 27-APR-1999 (first entry)

XX DE

DE HFE mutant (H63D-HFE) polypeptide sequence.

XX KW

KW HFE; beta-2-microglobulin; beta2m; iron overload; hemochromatosis;
transfusion; protein replacement therapy; hereditary hemochromatosis;
transferrin receptor; iron deficiency; anemia; mutant.

XX OS

OS Synthetic.

XX FH

FH Key Location/Qualifiers

FT FT

FT Misc-difference 2 /note= "indicated in the sequence listing as Arg"

FT FT

FT Misc-difference 41 /label= H63D

FT FT /note= "wild type His (of the mature protein sequence) is replaced by Asp"

FT FT

FT W09856814-A1.

PN PN

PN 17-DEC-1998.

PD PD

PD 12-JUN-1998; 98WO-US12436.

PF PF

PF 13-JUN-1997; 97US-0876010.

XX XX

PA (CALY) CALIFORNIA INST OF TECHNOLOGY.
 XX (PROG-) PROGENITOR INC.
 PI Bjorkman PJ, Feder JN, Schatzman RC;
 XX WPI; 1999-080886/07.
 DR
 XX
 PT New treatment of an iron overload disease - comprises use of HFE
 PT polypeptides provided in a complex with full length, wild type human
 PT (2m), useful in protein replacement therapy
 XX
 PS Claim 3; Page 14; 36pp; English.
 XX
 CC The present sequence represents a H63D-HFE mutant polypeptide. The HFE
 CC polypeptides (AA94295-297) provided in a complex with full length,
 CC wild type human beta-2-microglobulin (beta2m) form compositions in the
 CC treatment of primary iron overload diseases (e.g. haemochromatosis), or
 CC other iron overload conditions resulting from secondary causes (e.g.
 CC repeated transfusions). Data regarding the structure and function
 CC correlations of HFE polypeptides is useful in designing drugs that
 CC modulate the HFE gene and HFE activity. The polypeptides are also useful
 CC in protein replacement therapy for individuals possessing a defective
 CC HFE gene (e.g. Hereditary haemochromatosis). (Ant)agonists of the
 CC polypeptides are also useful in treating primary and secondary iron
 CC overload diseases. The modulators of the transferrin receptor are useful
 CC in treating iron deficiency conditions such as anemia, and in modulating
 CC the amount of iron transported into a cell. The HFE polypeptides provide
 CC a molecular basis for the relationship between HFE and iron metabolism,
 CC which enables treatment of iron overload and deficiency diseases.
 XX
 SQ Sequence 276 AA;
 Query Match 100.0%; Score 1520; DB 20; Length 276;
 Best Local Similarity 100.0%; Pred. No. 7.3e-135;
 Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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 Db 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDQLFVYDDERRRVEPTPWSSRISSQ 60
 Qy 61 MWLQSLQSLKGWDMFTVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
 Db 61 MWLQSLQSLKGWDMFTVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
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 Db 121 QDHLEFCPTDLWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
 Qy 181 DQVPPPLVKVTHHTVSSVTTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
 Db 181 DQVPPPLVKVTHHTVSSVTTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
 Qy 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 276
 Db 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 276
 RESULT 2
 ABG72686
 ID ABG72686 standard; protein; 276 AA.
 XX
 AC ABG72686;
 XX
 DT 05-MAR-2003 (first entry)
 XX
 DE Human haemochromatosis (HFE) mature protein, mutant H41D.
 XX
 KW Human; haemochromatosis; HFE; hereditary haemochromatosis;
 KW iron overload disease; iron deficiency disease; Beta2-microglobulin;
 KW Beta2m; transferrin receptor; anaemia; mutant; muten.
 XX
 OS Homo sapiens.
 OS Synthetic.

XX Key Location/Qualifiers
 FH Misc-difference 41
 FT /note= "wild-type His substituted by Asp"
 XX
 XX USG391852-B1.
 XX
 PD 21-MAY-2002.
 XX
 PF 12-JUN-1998; 98US-0094964.
 XX
 PR 13-JUN-1997; 97US-0876010.
 XX
 PA (BIRA) BIO-RAD LAB INC.
 PA (CALY) CALIFORNIA INST OF TECHNOLOGY.
 XX
 PI Feder JN, Bjorkman PJ, Schatzman RC;
 XX WPI; 2003-155377/15.
 DR
 XX Method of treating an iron overload disease comprises administration of
 PT a soluble complex comprising a 276 amino acid HFE polypeptide and a
 PT full length, wild-type human beta2m -
 XX
 PS Claim 2; Column 2; 17pp; English.
 XX
 CC The invention relates to a method of treating an iron overload disease
 CC comprising administration of a soluble complex comprising a 276 amino
 CC acid mature HFE (hereditary haemochromatosis gene protein) polypeptide
 CC (ABG72685-ABG72687) and a full length, wild-type human beta2m
 CC (beta2-microglobulin). In a HeLa cell based assay, binding and uptake of
 CC ¹²⁵I-transferrin in the presence of purified H63D-HFE/beta2m
 CC heterodimers was determined. At a concentration of 250 nM H63D-HFE/
 CC beta2m heterodimers, the transferrin receptor (TfR) displayed a KD for
 CC transferrin of 28 nM. At the same concentration of normal HFE/beta2m
 CC heterodimers, TfR displayed a KD for transferrin of 40 nM. In the absence
 CC of any HFE/beta2m heterodimers, TfR displayed a KD for transferrin of
 CC 7nM. It was observed that H63D-HFE/beta2m heterodimers were 30-40 % less
 CC efficient in decreasing TfR affinity for transferrin compared to
 CC wild-type HFE. The method is useful for treating iron overload diseases
 CC and iron deficiency e.g. anaemia. The present sequence is the H63D
 CC (residue 63 of the full length protein, 41 of the mature form)
 CC mutant form of mature HFE used to investigate the role of the His
 CC residue in transferrin receptor binding to transferrin.
 XX
 SQ Sequence 276 AA;
 Query Match 100.0%; Score 1520; DB 24; Length 276;
 Best Local Similarity 100.0%; Pred. No. 7.3e-135;
 Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
 Qy 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDQLFVYDDERRRVEPTPWSSRISSQ 60
 Db 1 RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDQLFVYDDERRRVEPTPWSSRISSQ 60
 Qy 61 MWLQSLQSLKGWDMFTVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
 Db 61 MWLQSLQSLKGWDMFTVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
 Qy 121 QDHLEFCPTDLWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
 Db 121 QDHLEFCPTDLWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
 Qy 181 DQVPPPLVKVTHHTVSSVTTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
 Db 181 DQVPPPLVKVTHHTVSSVTTLRCALNYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
 Qy 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 276
 Db 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 276
 RESULT 3

AA036871
ID AAB36871 standard; Protein; 348 AA.

XX AC AAB36871;

XX DT 21-FEB-2001 (first entry)

XX DE Human hereditary hemochromatosis 24d2 mutation protein.

XX KW HH; hereditary hemochromatosis; chelation agent;

XX KW T-cell differentiation factor; iron overload.

XX OS Homo sapiens;

XX PN US6140305-A.

XX PD 31-OCT-2000.

XX PF 04-APR-1997; 97US-0834497.

XX PR 04-APR-1996; 96US-0630912.

XX PR 16-APR-1996; 96US-0632673.

XX PR 23-MAY-1996; 96US-0652265.

XX PA (BIRA) BIO-RAD LAB INC.

XX PI Thomas-WJ, Drayna DT, Ghitke A, Ruddy D, Tsuchihashi Z, Wolff RK;

XX PI Feder JN;

XX DR WPI; 2001-006341/01.

XX DR N-PSDB; AAC68427.

XX PT New hereditary hemochromatosis gene products or polypeptides, useful
PT for treating hereditary hemochromatosis in a patient, and as a metal
PT chelation agent alleviating iron overload -

XX PS Claim 3; Fig 4; 108pp; English.

XX CC The present invention relates to hereditary hemochromatosis gene
CC products. These proteins may be used to treat a patient diagnosed as
CC having human hemochromatosis disease. It is also useful as a metal
CC chelation agent or as a T-cell differentiation factor, and for
CC alleviating iron overload. They may also be used in protein replacement
CC therapy for individuals having a defective human hemochromatosis gene.

XX SQ Sequence 348 AA;

Query Match 100.0%; Score 1520; DB 22; Length 348;

Best Local Similarity 100.0%; Pred. No. 9.9e-135;

Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RLIRSHSLHYLFMGASEQDLGLSLFALGYDDQLFVYDDERRRVEPTPWSSRISSQ 60

DB 23 RLIRSHSLHYLFMGASEQDLGLSLFALGYDDQLFVYDDERRRVEPTPWSSRISSQ 82

QY 61 MWLQSLQSLKGDHMTFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120

DB 83 MWLQSLQSLKGDHMTFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142

QY 121 QHLEFCPTDLWRAAEPRAPWTKLEWRHKIRARONRAYLERDCAQQLLELGRGVL 180

DB 143 QHLEFCPTDLWRAAEPRAPWTKLEWRHKIRARONRAYLERDCAQQLLELGRGVL 202

QY 181 DQOVPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEKDVLPNGDG 240

DB 203 DQOVPLVKVTHVTSVTLRCALNYPQNTMKWLKDKQPMDAKEPEKDVLPNGDG 262

QY 241 TYQGWITLAVPPGEQRYTCQVEHPGLDQPLIIVIE 276

DB 263 TYQGWITLAVPPGEQRYTCQVEHPGLDQPLIIVIE 298

RESULT 4

AAW94295

XX ID AAW94295 standard; peptide; 276 AA.

XX AC AAW94295;

XX DT 27-APR-1999 (first entry)

XX DE Wild-type HFE polypeptide sequence.

XX KW HFE; beta-2-microglobulin; beta2m; iron overload; hemochromatosis;
XX KW transfusion; protein replacement therapy; hereditary hemochromatosis;
XX KW transferrin receptor; iron deficiency; anemia.

XX OS Unidentified.

XX FH Key Location/Qualifiers

XX FT Misc-difference 2

XX FT /note= "indicated in the sequence listing as Arg"

XX PN W09856814-A1.

XX PD 17-DEC-1998.

XX PF 12-JUN-1998; 98WO-US12436.

XX PR 13-JUN-1997; 97US-0876010.

XX PA (CALY) CALIFORNIA INST OF TECHNOLOGY.

XX PA (PROG-) PROGENITOR INC.

XX PI Bjorkman PJ, Feder JN, Schatzman RC;

XX WPI; 1999-080886/07.

XX PT New treatment of an iron overload disease - comprises use of HFE

XX PT polypeptides provided in a complex with full length, wild type human

XX PT (2m), useful in protein replacement therapy

XX PS Claim 1; Page 13; 36pp; English.

XX CC The present sequence represents a wild-type HFE polypeptide. The HFE
XX polypeptides (AAW94295-297) provided in a complex with full length,
XX wild type human beta-2-microglobulin (beta2m) form compositions in the
XX treatment of primary iron overload diseases (e.g. hemochromatosis), or
XX other iron overload conditions resulting from secondary causes (e.g.
XX repeated transfusions). Data regarding the structure and function
XX correlations of HFE polypeptides is useful in designing drugs that
XX modulate the HFE gene and HFE activity. The polypeptides are also useful
XX in protein replacement therapy for individuals possessing a defective
XX HFE gene (e.g. Hereditary hemochromatosis). (Ant)agonists of the
XX polypeptides are also useful in treating primary and secondary iron
XX overload diseases. The modulators of the transferrin receptor are useful
XX in treating iron deficiency conditions such as anemia, and in modulating
XX the amount of iron transported into a cell. The HFE polypeptides provide
XX a molecular basis for the relationship between HFE and iron metabolism,
XX which enables treatment of iron overload and deficiency diseases.

XX SQ Sequence 276 AA;

Query Match 99.5%; Score 1513; DB 20; Length 276;

Best Local Similarity 99.6%; Pred. No. 3.3e-134;

Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

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DB 1 RLIRSHSLHYLFMGASEQDLGLSLFALGYDDQLFVYDDERRRVEPTPWSSRISSQ 60

QY 61 MWLQSLQSLKGDHMTFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120

DB 61 MWLQSLQSLKGDHMTFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120

QY 121 QHLEFCPTDLWRAAEPRAPWTKLEWRHKIRARONRAYLERDCAQQLLELGRGVL 180

Db 121 QDHFPCPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
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Db 181 DQOVPLVKVTHVTSVTLRCALNYPONTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
Db 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276

RESULT 5
ID ABG72685 standard; protein; 276 AA.
XX
AC ABG72685;
XX
DT 05-MAR-2003 (first entry)
XX
DE Human haemochromatosis (HFE) mature protein.
XX
KW Human; haemochromatosis; HFE; hereditary haemochromatosis;
KW iron overload disease; iron deficiency disease; Beta2-microglobulin;
KW Beta2m; transferrin receptor; anaemia.
XX
OS Homo sapiens.
XX
PN US6391852-B1.
XX
PD 21-MAY-2002.
XX
PF 12-JUN-1998; 98US-0094964.
XX
PR 13-JUN-1997; 97US-0876010.
XX
PA (BIRA) BIO-RAD LAB INC.
PA (CALY) CALIFORNIA INST OF TECHNOLOGY.
XX
PI Feder JN, Bjorkman PJ, Schatzman RC;
XX
DR WPI; 2003-155377/15.
XX
XX
PT Method of treating an iron overload disease comprises administration of
PT a soluble complex comprising a 276 amino acid HFE polypeptide and a
PT full length, wild-type human beta2m -
XX
PS Claim 1; Column 1; 17pp; English.
XX
XX
CC The invention relates to a method of treating an iron overload disease
CC comprising administration of a soluble complex comprising a 276 amino
CC acid mature HFE (hereditary haemochromatosis gene protein) polypeptide
CC (ABG72685-ABG72687) and a full length, wild-type human beta2m
CC (beta2-microglobulin). In a HeLa cell based assay, binding and uptake of
CC ⁵¹Fe-transferrin in the presence of purified H63D-HFE/beta2m
CC heterodimers was determined. At a concentration of 250 nM H63D-HFE/
CC beta2m heterodimers, the transferrin receptor (TfR) displayed a KD for
CC transferrin of 28 nM. At the same concentration of normal HFE/beta2m
CC heterodimers, TfR displayed a KD for transferrin of 40 nM. In the absence
CC of any HFE/beta2m heterodimers, TfR displayed a KD for transferrin of
CC 7nM. It was observed that H63D-HFE/beta2m heterodimers were 30-40 % less
CC efficient in decreasing TfR affinity for transferrin compared to
CC wild-type HFE. The method is useful for treating iron overload diseases
CC and iron deficiency e.g. anaemia. The present sequence is wild-type
CC mature HFE.
XX
SQ Sequence 276 AA;

Query Match 99.5%; Score 1513; DB 24; Length 276;
Best Local Similarity 99.6%; Pred. No. 3.3e-134;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRVPEPTPWSSRISSQ 60
|||||

Db 1 RLLRSHSLHYLFWGASEQDGLSLFEALGYVDDQLFVFDHESRRVPEPTPWSSRISSQ 60
QY 61 MWLQLSQSLSKGWDHMTVDFTWIMENHNHSKESHTLQVILGCEMOEDNSTEGYWKYGYDG 120
Db 61 MWLQLSQSLSKGWDHMTVDFTWIMENHNHSKESHTLQVILGCEMOEDNSTEGYWKYGYDG 120
QY 121 QDHFPCPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
Db 121 QDHFPCPTLDWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
QY 181 DQOVPLVKVTHVTSVTLRCALNYPONTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
Db 181 DQOVPLVKVTHVTSVTLRCALNYPONTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
Db 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276

RESULT 6
AAW36499
ID AAW36499 standard; Protein; 348 AA.
XX
AC AAW36499;
XX
DT 14-APR-1998 (first entry)
XX
DE Hereditary haemochromatosis gene product.
XX
KW Hereditary haemochromatosis; metal toxicity; diagnosis;
KW gene therapy; prenatal screening; human.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT Misc-difference 63 /note= "substituted by Asp in 24s2 mutant"
FT Misc-difference 65 /note= "substituted by Cys in 24d7 variant"
FT Misc-difference 282 /note= "substituted by Tyr in 24d1 mutant"
XX
PN WO9738137-A1.
XX
PD 16-OCT-1997.
XX
PF 04-APR-1997; 97WO-US06254.
XX
PR 23-MAY-1996; 96US-0652265.
PR 04-APR-1996; 96US-0630912.
PR 16-APR-1996; 96US-0632673.
XX
PA (MERC-) MERCATOR GENETICS INC.
PI Drayna DT, Feder JN, Gnirke A, Ruddy D, Thomas WJ;
PI Tsuchihashi Z, Wolff RK;
XX
DR WPI; 1997-512743/47.
DR N-PSDB; AAT96690, AAT96691.
XX
PT Hereditary haemochromatosis gene and variants - useful for diagnosis
PT and treatment of hereditary haemochromatosis disease
XX
PS Disclosure; Fig 4; 115pp; English.
XX
CC This polypeptide is the expression product of a novel human gene
CC (see AAT96690) whose mutated form is associated with hereditary
CC haemochromatosis (HH). A single mutation (24d1) in the HH gene
CC appears responsible for the majority of HH disease. This comprises
CC a G to A substitution that is present in 86% of affected
CC chromosomes and in 4% of unaffected chromosomes. It results in a
CC Cys to Tyr substitution in the encoded protein at a critical
CC disulphide bridge important for secondary structure. The following

CC are claimed: the 10825 bp genomic DNA sequence (1), a 1437 bp cDNA
CC sequence (1a) (see AAR96691) and their 24d1, 24d2 and 24d7 variants;
CC a cloning or expression vector; host cells; a peptide product
CC chosen from the HH gene product, its variants (24d1, 24d2 and
CC 24d7), or a peptide of at least 56 amino acid residues of these; an
CC antibody produced using the peptide as an immunogen; a method to
CC determine the presence or absence of the common HH gene mutation;
CC an animal model for the HH disease; metal chelation agents, T-cell
CC differentiation factors and therapeutic agents for the mitigation
CC of injury due to oxidative processes in vivo or mitigation of iron
CC overload; a method for screening potential therapeutic agents for
CC activity in connection with HH disease; an antisense oligonucleotide
CC directed against a transcriptional product of a nucleic acid
CC sequence as above; and oligonucleotides or pairs of oligonucleotides
CC covering a range of nucleotides from (1), (1a) or their variants,
CC useful for detecting a polymorphism in the HH gene. The invention
CC also relates to methods for screening for HH homozygotes, to HH
CC diagnosis, prenatal screening and diagnosis, and therapies of HH
CC disease, including gene therapy, protein- and antibody-based
CC therapeutics, and small molecule therapeutics.

XX
SQ Sequence 348 AA;
Query Match 99.5%; Score 1513; DB 18; Length 348;
Best Local Similarity 99.6%; Pred. No. 4.5e-134;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVYDDESRVPRTPWVSSRISSQ 60
DB 23 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVYDDESRVPRTPWVSSRISSQ 82
QY 61 MWLQSLQSLKGWDMFTVDFWTIMENHNHKSHTLQVILGCEQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGWDMFTVDFWTIMENHNHKSHTLQVILGCEQEDNSTEGYWKYGYDG 142
QY 121 QDHLFCPTDLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCEPAQLQQLLELGRGVL 180
DB 143 QDHLFCPTDLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCEPAQLQQLLELGRGVL 202
QY 181 DQOVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEKDPVLPNGDG 240
DB 203 DQOVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEKDPVLPNGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 298

RESULT 7
AAB19149
ID AAB19149 standard; Protein; 348 AA.
XX
AC AAB19149;
XX
DT 19-FEB-2001 (first entry)
XX
DE A human histocompatibility iron loading (HFE) protein.
KW Human; histocompatibility iron loading protein; HFE protein;
KW major histocompatibility complex; non-classical class I gene;
KW chromosome 6p; iron disorder; haemochromatosis.
XX
OS Homo sapiens.
XX
FH Key Location/Qualifiers
FT Peptide 1..22
FT /note= "signal peptide"
FT Misc-difference 63
FT /note= "when nucleotide 187 is mutated to G, then
FT this residue is Asp"
FT Misc-difference 65
FT /note= "when nucleotide 193 is mutated to T, then
FT this residue is Cys"

FT Domain 90..108
FT Misc-difference 93 /note= "alpha domain"
FT /note= "when nucleotide 277 is mutated to C, then
FT this residue is Arg"
FT Misc-difference 105 /note= "when nucleotide 314 is mutated to C, then
FT this residue is Thr"
WO200058515-A1.
05-OCT-2000.
24-MAR-2000; 2000WO-US07982.
26-MAR-1999; 99US-0277457.
(BILL-) BILLUPS-ROTHENBERG INC.
Rothenberg BE, Sawada-Hirai R, Barton JC;
WPI; 2000-647244/62.
N-PSDB; AAA96769.
Diagnosing an iron disorder e.g. hemochromatosis or a genetic
susceptibility to develop it, by determining the presence of a mutation
in exon 2 or an intron of a histocompatibility iron loading nucleic
acid -
XX
PS Disclosure; Page 3; 55pp; English.
XX
CC The present sequence represents a human histocompatibility iron loading
CC (HFE) protein. The HFE gene is a major histocompatibility (MHC)
CC non-classical class I gene located on chromosome 6p. Mutations in the
CC gene lead to iron disorders. The specification describes a method for
CC diagnosing an iron disorder or a genetic susceptibility to develop the
CC disorder in a mammal. The method comprises determining the presence of
CC a mutation in exon 2 or an intron of a HFE gene or protein. The mutation
CC is not a C to G missense mutation at nucleotide 187 of the sequence
CC given in A96769 (Genbank Accession number U60319). The presence of the
CC mutation indicates the disorder or the genetic susceptibility to the
CC disorder. The method is used to diagnose an iron disorder
CC e.g. haemochromatosis, or a genetic susceptibility to develop it.
XX
SQ Sequence 348 AA;
Query Match 99.5%; Score 1513; DB 21; Length 348;
Best Local Similarity 99.6%; Pred. No. 4.5e-134;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVYDDESRVPRTPWVSSRISSQ 60
DB 23 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVYDDESRVPRTPWVSSRISSQ 82
QY 61 MWLQSLQSLKGWDMFTVDFWTIMENHNHKSHTLQVILGCEQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGWDMFTVDFWTIMENHNHKSHTLQVILGCEQEDNSTEGYWKYGYDG 142
QY 121 QDHLFCPTDLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCEPAQLQQLLELGRGVL 180
DB 143 QDHLFCPTDLDWRAAEPRAPWTKLEWERHKIRARONRAYLERDCEPAQLQQLLELGRGVL 202
QY 181 DQOVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEKDPVLPNGDG 240
DB 203 DQOVPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEKDPVLPNGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 298
RESULT 8
AAB36869

ID AAB36869 standard; Protein; 348 AA.
XX
AC AAB36869;
XX
DT 21-FEB-2001 (first entry)
XX
DE Human hereditary hemochromatosis protein.
XX
KW HH; hereditary hemochromatosis; chelation agent;
KW T-cell differentiation factor; iron overload.
XX
OS Homo sapiens.
XX
PN US6140305-A.
XX
PD 31-OCT-2000.
XX
PF 04-APR-1997; 97US-0834497.
XX
PR 04-APR-1996; 96US-0630912.
PR 16-APR-1996; 96US-0632673.
PR 23-MAY-1996; 96US-0652265.
XX
PA (BIRA) BIO-RAD LAB INC.
XX
PI Thomas WJ, Drayna DT, Gnirke A, Ruddy D, Tsuchihashi Z, Wolff RK;
PI Feder JN;
XX
DR WPI; 2001-006341/01.
DR N-PSDB; AAC68425.
XX
PT New hereditary hemochromatosis gene products or polypeptides, useful
PT for treating hereditary hemochromatosis in a patient, and as a metal
PT chelation agent alleviating iron overload -
XX
XX Claim 1; Fig 4; 108pp; English.
XX
CC The present invention relates to hereditary hemochromatosis gene
CC products. These proteins may be used to treat a patient diagnosed as
CC having human hemochromatosis disease. It is also useful as a metal
CC chelation agent or as a T-cell differentiation factor, and for
CC alleviating iron overload. They may also be used in protein replacement
CC therapy for individuals having a defective human hemochromatosis gene.
XX
SQ Sequence 348 AA;
Query Match 99.5%; Score 1513; DB 22; Length 348;
Best Local Similarity 99.6%; Pred. No. 4.5e-134;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISQ 60
DB 23 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISQ 82
QY 61 MWLQLSQSLKGWDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGWDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDHLFCFDPDLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCAQQLLELGRGVL 180
DB 143 QDHLFCFDPDLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCAQQLLELGRGVL 202
QY 181 DQOVPLPVKVTHTVTSVTTLCRALNYYPQNTMKWLKDQPMDAKEFEKPDVLPNGDG 240
DB 203 DQOVPLPVKVTHTVTSVTTLCRALNYYPQNTMKWLKDQPMDAKEFEKPDVLPNGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 298

RESULT 9
AAB36872

ID AAB36872 standard; Protein; 348 AA.
XX
AC AAB36872;
XX
DT 21-FEB-2001 (first entry)
XX
DE Human hereditary hemochromatosis 24d1/2 mutation protein.
XX
KW HH; hereditary hemochromatosis; chelation agent;
KW T-cell differentiation factor; iron overload.
XX
OS Homo sapiens.
XX
PN US6140305-A.
XX
PD 31-OCT-2000.
XX
PF 04-APR-1997; 97US-0834497.
XX
PR 04-APR-1996; 96US-0630912.
PR 16-APR-1996; 96US-0632673.
PR 23-MAY-1996; 96US-0652265.
XX
PA (BIRA) BIO-RAD LAB INC.
XX
PI Thomas WJ, Drayna DT, Gnirke A, Ruddy D, Tsuchihashi Z, Wolff RK;
PI Feder JN;
XX
DR WPI; 2001-006341/01.
DR N-PSDB; AAC68428.
XX
PT New hereditary hemochromatosis gene products or polypeptides, useful
PT for treating hereditary hemochromatosis in a patient, and as a metal
PT chelation agent alleviating iron overload -
XX
XX Claim 4; Fig 4; 108pp; English.
XX
CC The present invention relates to hereditary hemochromatosis gene
CC products. These proteins may be used to treat a patient diagnosed as
CC having human hemochromatosis disease. It is also useful as a metal
CC chelation agent or as a T-cell differentiation factor, and for
CC alleviating iron overload. They may also be used in protein replacement
CC therapy for individuals having a defective human hemochromatosis gene.
XX
SQ Sequence 348 AA;
Query Match 99.3%; Score 1509; DB 22; Length 348;
Best Local Similarity 99.6%; Pred. No. 1.1e-133;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISQ 60
DB 23 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPTPWSSRISQ 82
QY 61 MWLQLSQSLKGWDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGWDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDHLFCFDPDLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCAQQLLELGRGVL 180
DB 143 QDHLFCFDPDLDWRAAEPRAPWTKLEWERHKIRARQNRAYLERDCAQQLLELGRGVL 202
QY 181 DQOVPLPVKVTHTVTSVTTLCRALNYYPQNTMKWLKDQPMDAKEFEKPDVLPNGDG 240
DB 203 DQOVPLPVKVTHTVTSVTTLCRALNYYPQNTMKWLKDQPMDAKEFEKPDVLPNGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 298

RESULT 10
AAU80035

Best Local Similarity 99.3%; Pred. No. 4.9e-133;
Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYDDQLFVYDDSRVPRTPWVSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASQDGLSLFEALGYDDQLFVYDDSRVPRTPWVSSRISSQ 82
QY 61 MWLQLSQSLKGWDHMTVDFTWIMENHNHKSHTLQVILGCEMOEDNSTEGWKYGYDG 120
DB 83 MWLQLSQSLKGWDHMTVDFTWIMENHNHKSHTLQVILGCEMOEDNSTEGWKYGYDG 142
QY 121 QDHLFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDPCPAQLQQLLELGRGVL 180
DB 143 QDHLFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDPCPAQLQQLLELGRGVL 202
QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMADAKEFPKDVLPNGDG 240
DB 203 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMADAKEFPKDVLPNGDG 262
QY 241 TYQGWITLAVPPGEGEORYTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPPGEGEORYTCQVEHPGLDQPLIVWE 298

RESULT 12

AAW94297
ID AAW94297 standard; peptide; 276 AA.
XX AAW94297;
XX
XX
XX 27-APR-1999 (first entry)
XX
DE HFE mutant (H111A/H145A-HFE) polypeptide sequence.
XX
KW HFE; beta-2-microglobulin; beta2m; iron overload; hemochromatosis;
KW transfusion; protein replacement therapy; hereditary hemochromatosis;
KW transferrin receptor; iron deficiency; anemia; mutant.
XX
OS Synthetic.
XX
XX
FH Key Location/Qualifiers
FT Misc-difference 2 /note= "indicated in the sequence listing as Arg"
FT
FT Misc-difference 89 /label= H111A
FT /note= "wild type His (of the mature protein sequence) is replaced by Ala"
FT
FT Misc-difference 123 /label= H145A
FT /note= "wild type His (of the mature protein sequence) is replaced by Ala"
FT
XX
XX WO9856814-A1.
XX
XX
XX 17-DEC-1998.
XX
XX 12-JUN-1998; 98WO-US12436.
XX
XX 13-JUN-1997; 97US-0876010.
XX
XX (CALY) CALIFORNIA INST OF TECHNOLOGY.
XX (PROG-) PROGENITOR INC.
XX
XX Bjorkman PJ., Feder JN, Schatzman RC;
XX
XX WPI; 1999-080886/07.
XX
XX New treatment of an iron overload disease - comprises use of HFE
XX polypeptides provided in a complex with full length, wild type human
XX (2m), useful in protein replacement therapy
XX
XX Claim 5; Page 15; 36pp; English.

CC The present sequence represents a H111A/H145A-HFE mutant polypeptide.
CC The HFE polypeptides (AAW94295-297) provided in a complex with full
CC length, wild type human beta-2-microglobulin (beta2m) form compositions
CC in the treatment of primary iron overload diseases (e.g.
CC hemochromatosis), or other iron overload conditions resulting from
CC secondary causes (e.g. repeated transfusions). Data regarding the
CC structure and function correlations of HFE polypeptides is useful in
CC designing drugs that modulate the HFE gene and HFE activity. The
CC polypeptides are also useful in protein replacement therapy for
CC individuals possessing a defective HFE gene (e.g. Hereditary
CC hemochromatosis). (Ant)agonists of the polypeptides are also useful in
CC treating primary and secondary iron overload diseases. The modulators of
CC the transferrin receptor are useful in treating iron deficiency
CC conditions such as anemia, and in modulating the amount of iron
CC transported into a cell. The HFE polypeptides provide a molecular basis
CC for the relationship between HFE and iron metabolism, which enables
CC treatment of iron overload and deficiency diseases.
XX
SQ Sequence 276 AA;

Query Match 98.2%; Score 1493; DB 20; Length 276;
Best Local Similarity 98.9%; Pred. No. 2.5e-132;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYDDQLFVYDDSRVPRTPWVSSRISSQ 60
DB 1 RLLRSHSLHYLFMGASQDGLSLFEALGYDDQLFVYDDSRVPRTPWVSSRISSQ 60
QY 61 MWLQLSQSLKGWDHMTVDFTWIMENHNHKSHTLQVILGCEMOEDNSTEGWKYGYDG 120
DB 61 MWLQLSQSLKGWDHMTVDFTWIMENHNHKSHTLQVILGCEMOEDNSTEGWKYGYDG 120
QY 121 QDHLFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDPCPAQLQQLLELGRGVL 180
DB 121 QDHLFCPTDLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDPCPAQLQQLLELGRGVL 180
QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMADAKEFPKDVLPNGDG 240
DB 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMADAKEFPKDVLPNGDG 240
QY 241 TYQGWITLAVPPGEGEORYTCQVEHPGLDQPLIVWE 276
DB 241 TYQGWITLAVPPGEGEORYTCQVEHPGLDQPLIVWE 276

RESULT 13

ABG72687
ID ABG72687 standard; protein; 276 AA.
XX
XX ABG72687;
XX
XX 05-MAR-2003 (first entry)
XX
XX Human haemochromatosis (HFE) mature protein, mutant H89A/H123A.
XX
XX Human; haemochromatosis; HFE; hereditary haemochromatosis;
KW iron overload disease; iron deficiency disease; Beta2-microglobulin;
KW Beta2m; transferrin receptor; anaemia; mutant; mutein.
XX
XX Homo sapiens.
OS Synthetic.
XX
XX Key Location/Qualifiers
FH Misc-difference 89 /note= "Wild-type His substituted by Ala"
FT
FT Misc-difference 123 /note= "Wild-type His substituted by Ala"
FT
XX US6391852-B1.
PN
XX 21-MAY-2002.
PD
XX
XX 12-JUN-1998; 98US-0094964.

XX PR 13-JUN-1997; 97US-0876010.
XX PA (BIRA) BIO-RAD LAB INC.
XX PA (CALY) CALIFORNIA INST OF TECHNOLOGY.
XX PI Feder JN, Bjorkman PJ, Schatzman RC;
XX WPI; 2003-155377/15.
XX
XX Method of treating an iron overload disease comprises administration of
PT a soluble complex comprising a 276 amino acid HFE polypeptide and a
PT full length, wild-type human beta2m -
XX
XX Claim 3; Column 2; 17pp; English.
XX
XX The invention relates to a method of treating an iron overload disease
CC comprising administration of a soluble complex comprising a 276 amino
CC acid mature HFE (hereditary haemochromatosis gene protein) polypeptide
CC (ABG72685-ABG72687) and a full length, wild-type human beta2m
CC (beta2-microglobulin). In a HeLa cell based assay, binding and uptake of
CC ⁵¹Fe-transferrin in the presence of purified H63D-HFE/beta2m
CC heterodimers was determined. At a concentration of 250 nM H63D-HFE/
CC beta2m heterodimers, the transferrin receptor (TfR) displayed a KD for
CC transferrin of 28 nM. At the same concentration of normal HFE/beta2m
CC heterodimers, TfR displayed a KD for transferrin of 40 nM. In the absence
CC of any HFE/beta2m heterodimers, TfR displayed a KD for transferrin of
CC 7nM. It was observed that H63D-HFE/beta2m heterodimers were 30-40 % less
CC efficient in decreasing TfR affinity for transferrin compared to
CC wild-type HFE. The method is useful for treating iron overload diseases
CC and iron deficiency e.g. anaemia. The present sequence is the H11A/H145A
CC (residues 111 and 145 of the full length protein, 89/123 of the mature
CC form) mutant from of mature HFE used to investigate the role of the His
CC residues in transferrin receptor binding to transferrin.
XX
XX SQ Sequence 276 AA;

Query Match 98.2%; Score 1493; DB 24; Length 276;
Best Local Similarity 98.9%; Pred. No. 2.5e-132;
Matches 273; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 1 RLLRSHLYLFWGASEODLGLSLFEALGVYDDQLFVYDDESRVPRTPWSSRISSQ 60
DB 1 RLLRSHLYLFWGASEODLGLSLFEALGVYDDQLFVYDDESRVPRTPWSSRISSQ 60
QY 61 MWLQSLQSGWDMFTVDFTWIMENHNSKESHTLQVILGCEMOEDNS-TEGWYKYGYDG 120
DB 61 MWLQSLQSGWDMFTVDFTWIMENHNSKESHTLQVILGCEMOEDNS-TEGWYKYGYDG 120
QY 121 QHLEFCPTLDWRAEPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 121 QDALEFCPTLDWRAEPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
QY 181 DQOVPLVAVKTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEFEFPKDVLPNGDG 240
DB 181 DQOVPLVAVKTHVTSSVTLRCALNYPQNTMKWLKDKQPMDAKEFEFPKDVLPNGDG 240
QY 241 TYQGWITLAVPPGEEQRYTCQVHPGLDQPLIWIWE 276
DB 241 TYQGWITLAVPPGEEQRYTCQVHPGLDQPLIWIWE 276

RESULT 14
AAB36873
ID AAB36873 standard; Protein; 361 AA.
XX
XX AAB36873;
XX
XX 21-FEB-2001 (first entry)
XX Rabbit leukocyte antigen.
XX
XX HH; hereditary hemochromatosis; chelation agent;
KW

KW T-cell differentiation factor; iron overload.
XX
XX Oryctolagus cuniculus.
XX
XX US6140305-A.
XX 31-OCT-2000.
XX
XX 04-APR-1997; 97US-0834497.
XX
XX 04-APR-1996; 96US-0630912.
XX 16-APR-1996; 96US-0632873.
XX 23-MAY-1996; 96US-0652265.
XX
XX (BIRA) BIO-RAD LAB INC.
XX
XX Thomas WJ, Drayna DT, Gnirke A, Ruddy D, Tsuchihashi Z, Wolff RK;
XX Feder JN;
XX WPI; 2001-006341/01.

XX New hereditary hemochromatosis gene products or polypeptides, useful
PT for treating hereditary hemochromatosis in a patient, and as a metal
PT chelation agent alleviating iron overload -
XX
XX Disclosure; Fig 7; 108pp; English.
XX
XX The present invention relates to hereditary hemochromatosis gene
CC products. These proteins may be used to treat a patient diagnosed as
CC having human hemochromatosis disease. It is also useful as a metal
CC chelation agent or as a T-cell differentiation factor, and for
CC alleviating iron overload. They may also be used in protein replacement
CC therapy for individuals having a defective human hemochromatosis gene.
XX
XX SQ Sequence 361 AA;

Query Match 34.4%; Score 523; DB 22; Length 361;
Best Local Similarity 40.1%; Pred. No. 9.1e-41;
Matches 111; Conservative 44; Mismatches 114; Indels 8; Gaps 7;
QY 5 SHSLHYLFWGASEODLGLSLFEALGVYDDQLFVYDDE--SRVPRTPWSSRISSQMW 62
DB 26 SHSMRYFTYSVRPGELGEPRTIIVGVYDDTQVFRDSDAASPRMEQRAFMW-GQVEPEY 84
QY 63 LQLSLSLQSGWDMFTVDFTWIMENHNSKE-SHTLQVILGCEMOEDNS-TEGWYKYGYDG 120
DB 85 DQQTQIAKTAQTAFRNLNTALRYNQSAAGSHFTQTFMGCEVWADGRFFHGYRAYDG 144
QY 121 QHLEFCPTLDWRAEPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 145 ADVIALNEDLRSWTAADTAQNTQKWEAAG-EAERHRAVLERECVEWLRRLVLEMGK 203
QY 181 DQOVPLVAVKTHVTSS-VTLRCALNYPQNTMKWLKDKQPMDAKEFEFPKDVLPNGD 239
DB 204 QRADPPKAVHTHPASDREATLRCWALGFYPAISLTWQDGDG-QTQDTLTVETRPGD 262
QY 240 GTYQGWITLAVPPGEEQRYTCQVHPGLDQPLIWIWE 276
DB 263 GTYQKVAWVPSGEEQRYTCVQHEGLPEPLTLTWE 299

RESULT 15
ABP68379
ID ABP68379 standard; Protein; 92 AA.
XX
XX ABP68379;
XX
XX 08-JAN-2003 (first entry)
XX Human colon specific protein, SEQ ID 120.
XX
XX Human; colon; cytostatic; vaccine; colon cancer; colon disorder;
KW metastasis.

XX	Homo sapiens.
OS	
XX	WO200277234-A2.
PN	
XX	
PD	03-OCT-2002.
XX	
XX	
PF	31-OCT-2001; 2001WO-US48414.
XX	
PR	31-OCT-2000; 2000US-244758P.
XX	
PA	(DIAD-) DIADEXUS INC.
XX	
PI	Sun Y, Recipon H, Ghosh MG, Liu C;
XX	
DR	WFI; 2003-018928/01.
XX	
PT	New isolated colon-specific nucleic acid molecule, useful for treating
PT	colon cancer, and diagnosing or monitoring the presence of metastases
PT	of colon cancer in a patient -
XX	
PS	Claim 11; Page 192; 216pp; English.
XX	
CC	The present invention relates to human colon specific nucleic acids
CC	(ABV93910-ABV94009) and proteins (ABP68360-ABP68435). The nucleic acids
CC	and proteins are useful for treating colon cancer and colon disorders,
CC	and diagnosing or monitoring the presence of colon disorders and
CC	metastases of colon cancer in a patient.
XX	
SQ	Sequence 92 AA;
	Query Match 33.8%; Score 514; DB 24; Length 92;
	Best Local Similarity 100.0%; Pred. No. 1e-40;
	Matches 92; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
QY	92 ESHTLQVLGCQEEDNSTGYWKYGVDGQDHLFCFDTLDWAABPRAWPTKLEWERHK 151
Db	1 ESHTLQVLGCQEEDNSTGYWKYGVDGQDHLFCFDTLDWAABPRAWPTKLEWERHK 60
QY	152 IRARONRAYLERDCPAQLQOLLEIGRGVLDQQ 183
Db	61 IRARONRAYLERDCPAQLQOLLEIGRGVLDQQ 92

Search completed: August 5, 2003, 13:08:23
Job time : 39 secs

GenCore version 5.1.6
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OM protein - protein search, using sw model

Run on: August 5, 2003, 13:07:04 ; Search time 14.5 seconds
(without alignments)
805.365 Million cell updates/sec

Title: US-10-092-404-2
Perfect score: 1520
Sequence: 1 RLLRSHSLHVLFWGASEQDL.....RYTCQVEHFGDQPLIVWE 276

Scoring table: BLOSUM62
Gapop 10.0 , Gapext 0.5

Searched: 328717 seqs, 42310858 residues

Total number of hits satisfying chosen parameters: 328717

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 45 summaries

Database : Issued Patents AA:*
1: /cgn2_6/ptodata/1/iaa/5A COMB.pap.*
2: /cgn2_6/ptodata/1/iaa/5B COMB.pap.*
3: /cgn2_6/ptodata/1/iaa/6A COMB.pap.*
4: /cgn2_6/ptodata/1/iaa/6B COMB.pap.*
5: /cgn2_6/ptodata/1/iaa/PCTUS COMB.pap.*
6: /cgn2_6/ptodata/1/iaa/backfile1.pap.*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	DB ID	Description
1	1520	100.0	276	4	US-09-094-964-2 Sequence 2, Appli
2	1520	100.0	348	3	US-08-552-265-6 Sequence 6, Appli
3	1520	100.0	348	3	US-08-834-497A-6 Sequence 6, Appli
4	1520	100.0	348	3	US-09-503-444A-6 Sequence 1, Appli
5	1513	99.5	276	4	US-09-094-964-1 Sequence 2, Appli
6	1513	99.5	348	3	US-08-552-265-2 Sequence 2, Appli
7	1513	99.5	348	3	US-08-834-497A-2 Sequence 2, Appli
8	1513	99.5	348	3	US-09-503-444A-2 Sequence 2, Appli
9	1513	99.5	348	4	US-09-277-457-2 Sequence 2, Appli
10	1513	99.5	348	4	US-09-679-729-2 Sequence 2, Appli
11	1509	99.3	348	3	US-08-552-265-8 Sequence 8, Appli
12	1509	99.3	348	3	US-08-834-497A-8 Sequence 8, Appli
13	1509	99.3	348	3	US-09-503-444A-8 Sequence 8, Appli
14	1502	98.8	348	3	US-08-552-265-4 Sequence 4, Appli
15	1502	98.8	348	3	US-08-834-497A-4 Sequence 4, Appli
16	1502	98.8	348	3	US-09-503-444A-4 Sequence 4, Appli
17	1493	98.2	276	4	US-09-094-964-3 Sequence 3, Appli
18	523	34.4	361	3	US-08-552-265-22 Sequence 22, Appli
19	523	34.4	361	3	US-08-834-497A-22 Sequence 22, Appli
20	523	34.4	361	3	US-09-503-444A-22 Sequence 22, Appli
21	517	34.0	364	4	US-08-514-372C-11 Sequence 11, Appli
22	514	33.8	365	3	US-08-552-265-23 Sequence 23, Appli
23	514	33.8	365	3	US-08-834-497A-23 Sequence 23, Appli
24	514	33.8	365	3	US-09-503-444A-23 Sequence 23, Appli
25	506	33.3	274	2	US-08-484-905-107 Sequence 107, App
26	506	33.3	274	3	US-08-481-985B-107 Sequence 107, App
27	506	33.3	274	3	US-08-370-476-107 Sequence 107, App

28	506	33.3	341	3	US-08-890-719-38 Sequence 38, Appli
29	505	33.2	365	2	US-08-484-905-97 Sequence 97, Appli
30	505	33.2	365	3	US-08-481-985B-97 Sequence 97, Appli
31	505	33.2	365	3	US-08-370-476-97 Sequence 97, Appli
32	504	33.2	274	2	US-08-484-905-108 Sequence 108, App
33	504	33.2	274	3	US-08-481-985B-108 Sequence 108, App
34	504	33.2	274	3	US-08-370-476-108 Sequence 108, App
35	504	33.2	365	2	US-08-484-905-100 Sequence 100, App
36	504	33.2	365	3	US-08-481-985B-100 Sequence 100, App
37	504	33.2	365	3	US-08-370-476-100 Sequence 100, App
38	503	33.1	274	1	US-08-222-851-1 Sequence 1, Appli
39	503	33.1	363	4	US-08-914-372C-37 Sequence 37, Appli
40	503	33.1	365	2	US-08-484-905-99 Sequence 99, Appli
41	503	33.1	365	3	US-08-481-985B-99 Sequence 99, Appli
42	503	33.1	365	3	US-08-370-476-99 Sequence 99, Appli
43	502	33.0	274	2	US-08-484-905-106 Sequence 106, App
44	502	33.0	274	3	US-08-481-985B-106 Sequence 106, App
45	502	33.0	274	3	US-08-370-476-106 Sequence 106, App

ALIGNMENTS

RESULT 1
US-09-094-964-2
; Sequence 2, Application US/09094964
; Patent No. 6391852
; GENERAL INFORMATION:
; APPLICANT: Feder, John N.
; APPLICANT: Bjorkman, Pamela J.
; APPLICANT: Schatzman, Randall C.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
; TITLE OF INVENTION: DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
; TITLE OF INVENTION: AND IRON DEFICIENCY DISEASES
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSES: Pennie & Edmonds, LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10036-2811
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: Windows
; SOFTWARE: FastSeq for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/094,964
; FILING DATE: June 12, 1998
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/876,010
; FILING DATE: June 13, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0074-999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-493-4935
; TELEFAX: 650-493-5556
; TELEX: 66141 PENNIE
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; TYPE: amino acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; MOLECULE TYPE: peptide
US-09-094-964-2

Query Match 100.0%; Score 1520; DB 4; Length 276;
Best Local Similarity 100.0%; Pred. No. 2.2e-142;

Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDSSRRVPRTPWSSRISQ 60
1 RLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDSSRRVPRTPWSSRISQ 60
DB 1 RLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDSSRRVPRTPWSSRISQ 60
QY 61 MWLQLSQSLSKGDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
61 MWLQLSQSLSKGDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 121 QDHLFCPDTLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
121 QDHLFCPDTLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
DB 121 QDHLFCPDTLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
QY 181 DQOVPLVAVKTHVTSSVTLRCALNYPONITMKWLKDKQPMDAKEPEPKDVLNGDG 240
181 DQOVPLVAVKTHVTSSVTLRCALNYPONITMKWLKDKQPMDAKEPEPKDVLNGDG 240
DB 181 DQOVPLVAVKTHVTSSVTLRCALNYPONITMKWLKDKQPMDAKEPEPKDVLNGDG 240
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
DB 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276

RESULT 2

US-08-652-265-6
; Sequence 6, Application US/08652265
; Patent No. 6025130
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: Hereditary Hemochromatosis Gene
; NUMBER OF SEQUENCES: 44
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew LLP
; STREET: Two Embarcadero Center, Eighth Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/652,265
; FILING DATE: 23-MAY-1996
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M.
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 17957-000500
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 576-0200
; TELEFAX: (415) 576-0300
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-652-265-6

Query Match 100.0%; Score 1520; DB 3; Length 348;
Best Local Similarity 100.0%; Pred. No. 3e-142;
Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDSSRRVPRTPWSSRISQ 60

DB 23 RLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDSSRRVPRTPWSSRISQ 82
QY 61 MWLQLSQSLSKGDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLSKGDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDHLFCPDTLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLFCPDTLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQOVPLVAVKTHVTSSVTLRCALNYPONITMKWLKDKQPMDAKEPEPKDVLNGDG 240
DB 203 DQOVPLVAVKTHVTSSVTLRCALNYPONITMKWLKDKQPMDAKEPEPKDVLNGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIWIWE 298

RESULT 3

US-08-834-497A-6
; Sequence 6, Application US/08834497A
; Patent No. 6149305
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: HEREDITARY HEMOCHROMATOSIS GENE PRODUCTS
; NUMBER OF SEQUENCES: 76
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036-2811
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: FastSeq for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/834,497A
; FILING DATE: 04-APR-1997
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/652,265
; FILING DATE: 23-MAY-1996
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/632,673
; FILING DATE: 16-APR-1996
; CLASSIFICATION: 514
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/630,912
; FILING DATE: 04-APR-1996
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M.
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0056-999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-493-4935
; TELEFAX: 650-493-5556
; TELEX: 66141 PENNIE
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids

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; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-834-497A-6

Query Match      100.0%; Score 1520; DB 3; Length 348;
Best Local Similarity 100.0%; Pred. No. 3e-142;
Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDLGLSLFEALGYVDQLFVYDDERRRVEPRTPWSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASQDLGLSLFEALGYVDQLFVYDDERRRVEPRTPWSSRISSQ 82
QY 61 MWLQSLQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDHLEFCPTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLEFCPTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
DB 203 DQVPPPLVKVTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 298

RESULT 4
US-09-503-444A-6
; Sequence 6, Application US/09503444A
; Patent No. 6228594
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: Hereditary Hemochromatosis Gene
; NUMBER OF SEQUENCES: 44
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: Wordperfect Version 8
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/503,444A
; FILING DATE: 14-Feb-2000
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/652,265
; FILING DATE: 23-May-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/632,673
; FILING DATE: 16-Apr-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/630,912
; FILING DATE: 04-Apr-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M.
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0088-999
```

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; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 212-790-9090
; TELEFAX: 212-869-9741
; TELEX: 66141
; INFORMATION FOR SEQ ID NO: 6:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-09-503-444A-6

Query Match      100.0%; Score 1520; DB 3; Length 348;
Best Local Similarity 100.0%; Pred. No. 3e-142;
Matches 276; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDLGLSLFEALGYVDQLFVYDDERRRVEPRTPWSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASQDLGLSLFEALGYVDQLFVYDDERRRVEPRTPWSSRISSQ 82
QY 61 MWLQSLQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLQSLKGDHMFVDFWTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDHLEFCPTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLEFCPTLDWRAAEPRAPWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
DB 203 DQVPPPLVKVTHVTSSVTLRCRALNYYPQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVIE 298

RESULT 5
US-09-094-964-1
; Sequence 1, Application US/09094964
; Patent No. 6391852
; GENERAL INFORMATION:
; APPLICANT: Feder, John N.
; APPLICANT: Bjorkman, Pamela J.
; APPLICANT: Schatzman, Randall C.
; TITLE OF INVENTION: METHODS AND COMPOSITIONS FOR
; TITLE OF INVENTION: DIAGNOSIS AND TREATMENT OF IRON OVERLOAD DISEASES
; NUMBER OF SEQUENCES: 5
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds, LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: NY
; COUNTRY: USA
; ZIP: 10036-2811
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: Windows
; SOFTWARE: FastSeq for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/094,964
; FILING DATE: June 12, 1998
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/876,010
; FILING DATE: June 13, 1997
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M.
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0074-999
; TELECOMMUNICATION INFORMATION:
```

TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 1:
SEQUENCE CHARACTERISTICS:
LENGTH: 276 amino acids
TYPE: amino acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: peptide
US-09-094-964-1

Query Match 99.5%; Score 1513; DB 4; Length 276;
Best Local Similarity 99.6%; Pred. No. 1.1e-141;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQLFVYDDERRRVEPRTPWSSRISSQ 60
DB 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQLFVYDDERRRVEPRTPWSSRISSQ 60
QY 61 MWLQSLKGDHMTVDFTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 61 MWLQSLKGDHMTVDFTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
QY 121 QDHLFCFDDTLDRAAEPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
DB 121 QDHLFCFDDTLDRAAEPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
DB 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
DB 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276

RESULT 6
US-08-652-265-2
Sequence 2, Application US/08652265
Patent No. 6025130
GENERAL INFORMATION:
APPLICANT: Thomas, Winston J.
APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: Hereditary Hemochromatosis Gene
NUMBER OF SEQUENCES: 44
CORRESPONDENCE ADDRESS:
ADDRESSEE: Townsend and Townsend and Crew LLP
STREET: Two Embarcadero Center, Eighth Floor
CITY: San Francisco
STATE: California
COUNTRY: USA
ZIP: 94111-3834
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: PatentIn Release #1.0, Version #1.30
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/652,265
FILING DATE: 23-MAY-1996
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Smith, William M.
REGISTRATION NUMBER: 30,223
REFERENCE/DOCKET NUMBER: 17957-000500
TELECOMMUNICATION INFORMATION:
TELEPHONE: (415) 576-0200

TELEFAX: (415) 576-0300
INFORMATION FOR SEQ ID NO: 2:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-652-265-2

Query Match 99.5%; Score 1513; DB 3; Length 348;
Best Local Similarity 99.6%; Pred. No. 1.1e-141;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQLFVYDDERRRVEPRTPWSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASQDGLSLFEALGYVDQLFVYDDERRRVEPRTPWSSRISSQ 82
QY 61 MWLQSLKGDHMTVDFTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQSLKGDHMTVDFTIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDHLFCFDDTLDRAAEPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLFCFDDTLDRAAEPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 240
DB 203 DQVPPPLVKVTHVTSSVTLRCRALNYPQNTMKWLKDKQPMDAKEPEPKDVLPGDG 262
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 298

RESULT 7
US-08-834-497A-2
Sequence 2, Application US/08834497A
Patent No. 6140305
GENERAL INFORMATION:
APPLICANT: Thomas, Winston J.
APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: HEREDITARY HEMOCHROMATOSIS GENE PRODUCTS
NUMBER OF SEQUENCES: 76
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036-2811
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: FastSeq for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/834,497A
FILING DATE: 04-APR-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/652,265
FILING DATE: 23-MAY-1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/632,673
FILING DATE: 16-APR-1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:

```

; APPLICATION NUMBER: US 08/630,912
; FILING DATE: 04-APR-1996
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M.
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0056-999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 650-493-4935
; TELEFAX: 650-493-5556
; TELEX: 66141 PENNIE
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-08-834-497A-2

Query Match 99.5%; Score 1513; DB 3; Length 348;
Best Local Similarity 99.6%; Pred. No. 1.5e-141;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVFDHESRRVEPRTPWVSSRISSQ 60
DB 23 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVFDHESRRVEPRTPWVSSRISSQ 82
QY 61 MWLQLSQSLKGDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDHLFCPTDLWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLFCPTDLWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQVPLVAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
DB 203 DQVPLVAVPPGEEQRYTCQVEHPGLDQPLIVWE 298
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 298

RESULT 8
US-09-503-444A-2
; Sequence 2, Application US/09503444A
; Patent No. 6228594
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: Hereditary Hemochromatosis Gene
; NUMBER OF SEQUENCES: 44
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Pennie & Edmonds LLP
; STREET: 1155 Avenue of the Americas
; CITY: New York
; STATE: New York
; COUNTRY: USA
; ZIP: 10036
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: WordPerfect Version 8
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/503,444A
; FILING DATE: 14-Feb-2000

; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/652,265
; FILING DATE: 23-May-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/632,673
; FILING DATE: 16-Apr-1996
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/630,912
; FILING DATE: 04-Apr-1996
; ATTORNEY/AGENT INFORMATION:
; NAME: Poissant, Brian M.
; REGISTRATION NUMBER: 28,462
; REFERENCE/DOCKET NUMBER: 8907-0088-999
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 212-790-9090
; TELEFAX: 212-869-9741
; TELEX: 66141
; INFORMATION FOR SEQ ID NO: 2:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
; US-09-503-444A-2

Query Match 99.5%; Score 1513; DB 3; Length 348;
Best Local Similarity 99.6%; Pred. No. 1.5e-141;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

QY 1 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVFDHESRRVEPRTPWVSSRISSQ 60
DB 23 RLLRSHSLHYLFWGASEQDLGLSLFEALGYVDDQLFVFDHESRRVEPRTPWVSSRISSQ 82
QY 61 MWLQLSQSLKGDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGDHMTVDFTWIMENHNHSHKESHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDHLFCPTDLWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLFCPTDLWRAAPRAWPTKLEWERHKIRARQNRAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQVPLVAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
DB 203 DQVPLVAVPPGEEQRYTCQVEHPGLDQPLIVWE 298
QY 241 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 276
DB 263 TYQGWITLAVPPGEEQRYTCQVEHPGLDQPLIVWE 298

RESULT 9
US-09-277-457-2
; Sequence 2, Application US/09277457
; Patent No. 6355425
; GENERAL INFORMATION:
; APPLICANT: Rothenberg, Barry E.
; APPLICANT: Sawada-Hirai, Ritsuko
; APPLICANT: Barton, James C.
; TITLE OF INVENTION: MUTATIONS ASSOCIATED WITH IRON DISORDERS
; FILE REFERENCE: 10653/002001
; CURRENT APPLICATION NUMBER: US/09/277,457
; CURRENT FILING DATE: 1999-03-26
; NUMBER OF SEQ ID NOS: 30
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 348
; TYPE: PRT
; ORGANISM: Homo Sapiens
; US-09-277-457-2

Query Match 99.5%; Score 1513; DB 4; Length 348;
Best Local Similarity 99.6%; Pred. No. 1.5e-141;
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Matches	275;	Conservative	0;	Mismatches	1;	Indels	0;	Gaps	0;
Qy	1	RLLRSHSLHYLFMGASQDGLSLFEALGYVDQLFVYFDDSRRLVEPRTPWVSSRISSQ	60						
Db	23	RLLRSHSLHYLFMGASEQDLGLSLFEALGYVDQLFVYFDDSRRLVEPRTPWVSSRISSQ	82						
Qy	61	MWLQSLQSLKGWDHMTVDFTWMENHNHSHKESHTLQVILGCEMQRDNSTEGYWKYGDG	120						
Db	83	MWLQSLQSLKGWDHMTVDFTWMENHNHSHKESHTLQVILGCEMQRDNSTEGYWKYGDG	142						
Qy	121	QDHLFCPDTLLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL	180						
Db	143	QDHLFCPDTLLDWRAAPRAWPTKLEWERHKIRARONRAYLERDCPAQLQQLLELGRGVL	202						
Qy	181	DOQVEPLVKVTHHYTSSVTTILRCALNYFPONTIMKWLKDQKPMADKEPEPKDVLPNGDG	240						
Db	203	DOQVEPLVKVTHHYTSSVTTILRCALNYFPONTIMKWLKDQKPMADKEPEPKDVLPNGDG	262						
Qy	241	TVQWGMITLAVPPGEEQRYTCQVEHPGLDQPLIUIVE	276						
Db	263	TVQWGMITLAVPPGEEQRYTCQVEHPGLDQPLIUIVE	298						

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RESULT 10
US-09-679-729-2
; Sequence 2, Application US/09679729
; Patent No. 6509442
; GENERAL INFORMATION:
; APPLICANT: Rothenberg, Barry E.
; APPLICANT: Sawada-Hirai, Ritsuko
; APPLICANT: Barcon, James C.
; TITLE OF INVENTION: MUTATIONS ASSOCIATED WITH IRON DISORDERS
; FILE REFERENCE: 24065-004 DIV
; CURRENT APPLICATION NUMBER: US/09/679,729
; CURRENT FILING DATE: 2000-10-04
; PRIOR APPLICATION NUMBER: 09/277,457
; PRIOR FILING DATE: 1999-03-26
; NUMBER OF SEQ ID NOS: 30
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 2
; LENGTH: 348
; TYPE: PRT
; ORGANISM: Homo Sapiens
US-09-679-729-2

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Query Match	99.5%;	Score 1513;	DB 4;	Length 348;
Best Local Similarity	99.6%;	Pred. No. 1.5e-141;		
Matches 275;	Conservative 0;	Mismatches 1;	Indels 0;	Gaps 0;
QY	1	RLLRSHSLHYLFMGASQDGLGLSLFEALGYVDDQLFVYDDSRVRVEPTPWVGSRISSQ	60	
Db	23	RLLRSHSLHYLFMGASBQDGLGLSLFEALGYVDDQLFVYDDSRVRVEPTPWVGSRISSQ	82	
QY	61	MWLQLSQSLKGWDHMFVDFWTIMENHNHNSKESHTLQVILGCMEQEDNSTGYWKYGYDG	120	
Db	83	MWLQLSQSLKGWDHMFVDFWTIMENHNHNSKESHTLQVILGCMEQEDNSTGYWKYGYDG	142	
QY	121	QDHLEFCDDTLWDRAEAPRAWPTKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGVL	180	
Db	143	QDHLEFCDDTLWDRAEAPRAWPTKLEWERHKIRARONRAYLERDPCPAQLQQLLELGRGVL	202	
QY	181	DQVPPPLVKVTHVHTSSVTLRLCRALNYYPQNTTMKWLKDQKPMDAKEFEKPDVLPNGDG	240	
Db	203	DQVPPPLVKVTHVHTSSVTLRLCRALNYYPQNTTMKWLKDQKPMDAKEFEKPDVLPNGDG	262	
QY	241	TYQGWITLAVPPGEEQRYTCQVHPGLDQPLIWIIE	276	
Db	263	TYQGWITLAVPPGEEQRYTCQVHPGLDQPLIWIIE	298	

RESULT 11
US-08-652-265-8
; Sequence 8, Application US/086522265

```

; Patent No. 605130
;
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
;
; TITLE OF INVENTION: Hereditary Hemochromatosis Gene
;
; NUMBER OF SEQUENCES: 44
;
; CORRESPONDENCE ADDRESSES:
; ADDRESSEE: Townsend and Townsend and Crew LLP
; STREET: Two Embarcadero Center, Eighth Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834
;
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.30
;
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/652,265
; FILING DATE: 23-MAY-1996
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M.
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 17957-000500
;
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 576-0200
; TELEFAX: (415) 576-0300
;
; INFORMATION FOR SEQ ID NO: 8:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
;
; US-08-652-265-8
;
; Query Match 99.3%; Score 1509; DB 3; Length
; Best Local Similarity 99.6%; Pred. No. 3.6e-141; Ind.
; Matches 275; Conservative 0; Mismatches 1; Indels 0
;
; QY 1 RLLRSHSLHYLFPMGASEODLGLSFALGYVDDQLFVYDDDESE
;
; DB 23 RLLRSHSLHYLFPMGASEODLGLSFALGYVDDQLFVYDDDESE
;
; QY 61 MWLQLSQSLKGDHMTFTVDFWTIMENHNHNSKESHTLQVILGCEN
;
; DB 83 MWLQLSQSLKGDHMTFTVDFWTIMENHNHNSKESHTLQVILGCEN
;
; QY 121 QDHLFECDDTLDWRAAEPRAVPTKLEWRHKIRARQNRAVYLEIR
;
; DB 143 QDHLFECDDTLDWRAAEPRAVPTKLEWRHKIRARQNRAVYLEIR
;
; QY 181 DQOVPEPLVKVTHVHTVSSVTTLRCRALNYYPONIIMKWLKDQKQPN
;
; DB 203 DQOVPEPLVKVTHVHTVSSVTTLRCRALNYYPONIIMKWLKDQKQPN
;
; QY 241 TYQGMITLAVPPGEQRVYTCVQEHPLDQPLVIWE 276
;
; DB 263 TYQGMITLAVPPGEQRVYTCVQEHPLDQPLVIWE 298

```

RESULT 12
US-08-834-497A-8
; Sequence 8, Application US/08834497A
; Patent No. 6140305
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.

APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: HEREDITARY HEMOCHROMATOSIS GENE PRODUCTS
NUMBER OF SEQUENCES: 76
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036-2811
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: FASTSQ for Windows Version 2.0b
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/834,497A
FILING DATE: 04-APR-1997
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/652,265
FILING DATE: 23-MAY-1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/632,673
FILING DATE: 16-APR-1996
CLASSIFICATION: 514
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/630,912
FILING DATE: 04-APR-1996
CLASSIFICATION: 514
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M.
REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0056-999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 650-493-4935
TELEFAX: 650-493-5556
TELEX: 66141 PENNIE
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-08-834-497A-8

Query Match 99.3%; Score 1509; DB 3; Length 348;
Best Local Similarity 99.6%; Pred. No. 3.6e-141;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDSSRRVPEPTPWSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDSSRRVPEPTPWSSRISSQ 82
QY 61 MWLQLSQSLKGDHMTFTVDFWTIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGDHMTFTVDFWTIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDHLFCPTDLWRAAEPRAPWTKLEWERHKIRARQNAYLERDCPAQLQQLLELGRGVL 180
DB 143 QDHLFCPTDLWRAAEPRAPWTKLEWERHKIRARQNAYLERDCPAQLQQLLELGRGVL 202
QY 181 DQQVPLVKVTHVTSVTLRCAALNYYPONTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
DB 203 DQQVPLVKVTHVTSVTLRCAALNYYPONTMKWLKDKQPMDAKEPEPKDVLNPGDG 262
QY 241 TYQGWITLAVPPGEQRYTCQVEHPGLDQPLIWIWE 276

DB 263 TYQGWITLAVPPGEQRYTCQVEHPGLDQPLIWIWE 298
RESULT 13
US-09-503-444A-8
Sequence 8, Application US/09503444A
Patent No. 6228594
GENERAL INFORMATION:
APPLICANT: Thomas, Winston J.
APPLICANT: Drayna, Dennis T.
APPLICANT: Feder, John N.
APPLICANT: Gnirke, Andreas
APPLICANT: Ruddy, David
APPLICANT: Tsuchihashi, Zenta
APPLICANT: Wolff, Roger K.
TITLE OF INVENTION: Hereditary Hemochromatosis Gene
NUMBER OF SEQUENCES: 44
CORRESPONDENCE ADDRESS:
ADDRESSEE: Pennie & Edmonds LLP
STREET: 1155 Avenue of the Americas
CITY: New York
STATE: New York
COUNTRY: USA
ZIP: 10036
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: Windows 95
SOFTWARE: WordPerfect Version 8
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/09/503,444A
FILING DATE: 14-Feb-2000
CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/652,265
FILING DATE: 23-May-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/632,673
FILING DATE: 16-Apr-1996
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/630,912
FILING DATE: 04-Apr-1996
ATTORNEY/AGENT INFORMATION:
NAME: Poissant, Brian M.
REGISTRATION NUMBER: 28,462
REFERENCE/DOCKET NUMBER: 8907-0088-999
TELECOMMUNICATION INFORMATION:
TELEPHONE: 212-790-9090
TELEFAX: 212-869-9741
TELEX: 66141
INFORMATION FOR SEQ ID NO: 8:
SEQUENCE CHARACTERISTICS:
LENGTH: 348 amino acids
TYPE: amino acid
TOPOLOGY: linear
MOLECULE TYPE: protein
US-09-503-444A-8

Query Match 99.3%; Score 1509; DB 3; Length 348;
Best Local Similarity 99.6%; Pred. No. 3.6e-141;
Matches 275; Conservative 0; Mismatches 1; Indels 0; Gaps 0;
QY 1 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDSSRRVPEPTPWSSRISSQ 60
DB 23 RLLRSHSLHYLFMGASEQDGLSLFEALGYVDDQLFVYDDSSRRVPEPTPWSSRISSQ 82
QY 61 MWLQLSQSLKGDHMTFTVDFWTIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDG 120
DB 83 MWLQLSQSLKGDHMTFTVDFWTIMENHNHKSHTLQVILGCEMQEDNSTEGYWKYGYDG 142
QY 121 QDHLFCPTDLWRAAEPRAPWTKLEWERHKIRARQNAYLERDCPAQLQQLLELGRGVL 180

Db 143 QDHLFCFDDTLDRABPRAWPTKLEWERHKBIRARONRAYLERDCPAQLQQLLELGRGVL 202
Qy 181 DQOVPLVAVKTHVTSSVTLRCALNYYQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240
Db 203 DQOVPLVAVKTHVTSSVTLRCALNYYQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262
Qy 241 TYQGWITLAVPPGEEQRYTCQVEHGLDQPLIVWE 276
Db 263 TYQGWITLAVPPGEEQRYTCQVEHGLDQPLIVWE 298

RESULT 14
US-08-652-265-4
; Sequence 4, Application US/08652265
; Patent No. 6025130
; GENERAL INFORMATION:
; APPLICANT: Thomas, Winston J.
; APPLICANT: Drayna, Dennis T.
; APPLICANT: Feder, John N.
; APPLICANT: Gnirke, Andreas
; APPLICANT: Ruddy, David
; APPLICANT: Tsuchihashi, Zenta
; APPLICANT: Wolff, Roger K.
; TITLE OF INVENTION: Hereditary Hemochromatosis Gene
; NUMBER OF SEQUENCES: 44
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Townsend and Townsend and Crew LLP
; STREET: Two Embarcadero Center, Eighth Floor
; CITY: San Francisco
; STATE: California
; COUNTRY: USA
; ZIP: 94111-3834
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/652,265
; FILING DATE: 23-MAY-1996
; CLASSIFICATION: 514
; ATTORNEY/AGENT INFORMATION:
; NAME: Smith, William M.
; REGISTRATION NUMBER: 30,223
; REFERENCE/DOCKET NUMBER: 17957-000500
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (415) 576-0200
; TELEFAX: (415) 576-0300
; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-652-265-4

Query Match 98.8%; Score 1502; DB 3; Length 348;
Best Local Similarity 99.3%; Pred. No. 1.8e-140;
Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 1 RLLRSHSLHYLFGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWVSSRISSQ 60
Db 23 RLLRSHSLHYLFGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWVSSRISSQ 82
Qy 61 MWLQSLKSGWDMFTVDFTWMENHNHSHKESHTLQVILGCMQEDNSTEGYWKYGYDG 120
Db 83 MWLQSLKSGWDMFTVDFTWMENHNHSHKESHTLQVILGCMQEDNSTEGYWKYGYDG 142
Qy 121 QDHLFCFDDTLDRABPRAWPTKLEWERHKBIRARONRAYLERDCPAQLQQLLELGRGVL 180
Db 143 QDHLFCFDDTLDRABPRAWPTKLEWERHKBIRARONRAYLERDCPAQLQQLLELGRGVL 202
Qy 181 DQOVPLVAVKTHVTSSVTLRCALNYYQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 240

Db 203 DQOVPLVAVKTHVTSSVTLRCALNYYQNTMKWLKDKQPMDAKEPEPKDVLNPGDG 262
Qy 241 TYQGWITLAVPPGEEQRYTCQVEHGLDQPLIVWE 276
Db 263 TYQGWITLAVPPGEEQRYTCQVEHGLDQPLIVWE 298

RESULT 15
US-08-834-497A-4
; Sequence 4, Application US/08834497A
; Patent No. 6140305
; GENERAL INFORMATION:
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; TITLE OF INVENTION: HEREDITARY HEMOCHROMATOSIS GENE PRODUCTS
; NUMBER OF SEQUENCES: 76
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; STREET: 1155 Avenue of the Americas
; CITY: New York
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; COUNTRY: USA
; ZIP: 10036-2811
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: Windows 95
; SOFTWARE: FastSEQ for Windows Version 2.0b
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/834,497A
; FILING DATE: 04-APR-1997
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; APPLICATION NUMBER: US 08/652,265
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; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/632,673
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; CLASSIFICATION: 514
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; INFORMATION FOR SEQ ID NO: 4:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 348 amino acids
; TYPE: amino acid
; TOPOLOGY: linear
; MOLECULE TYPE: protein
US-08-834-497A-4

Query Match 98.8%; Score 1502; DB 3; Length 348;
Best Local Similarity 99.3%; Pred. No. 1.8e-140;
Matches 274; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
Qy 1 RLLRSHSLHYLFGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWVSSRISSQ 60
Db 23 RLLRSHSLHYLFGASEQDGLSLFEALGYVDDQLFVYDDERRRVEPRTPWVSSRISSQ 82

Qy	61	MWLSQSLKGDHMFVDFWTFIMENNNHSHKESHTLQVILGCENQEDNSTEGYWKYGYDG	120
Db	83	MWLSQSLKGDHMFVDFWTFIMENNNHSHKESHTLQVILGCENQEDNSTEGYWKYGYDG	142
Qy	121	QDHLFCPCDPTLDWEAEPRAWPTKLEWERHKIRARONRAYLERDCPAQLOQLLELGRGVL	180
Db	143	QDHLFCPCDPTLDWEAEPRAWPTKLEWERHKIRARONRAYLERDCPAQLOQLLELGRGVL	202
Qy	181	DQVPPPLVKVTHHTVSSVTTLRCALNYYPNITMKWLKDKQPMDAKEFEFPKDVLPNGDG	240
Db	203	DQVPPPLVKVTHHTVSSVTTLRCALNYYPNITMKWLKDKQPMDAKEFEFPKDVLPNGDG	262
Qy	241	TYQGWITLAVPPGGEQRYTCQVEHPGLDQPLIVIWE	276
Db	263	TYQGWITLAVPPGGEQRYTCQVEHPGLDQPLIVIWE	298

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